MEMBER ROSEN: But that below 100 microns, 1 and we have reasonable assurance of that. That's 2 pedal to the metal all the way up to however many 3 qigawatt days per ton I want, right? 4 CHAIRMAN POWERS: Ten to the sixth, as a 5 matter of fact. 6 MR. MONTGOMERY: It's a straight line 7 after this. 8 It's a straight line, CHAIRMAN POWERS: 9 not because of what the fuel is doing, but because of 10 his capping the outside parameters. 11 MR. MONTGOMERY: Well, inherent in this 12 there's a burnup effect coming from the fuel pellet, 13 but the cladding ductility saturates and that's the 14 reason that the PCMI loading still remains the same. 15 And it's fairly asymptotic, yes. 16 MEMBER ROSEN: This is crucial. 17 what this work is saying is that as long as you keep 18 oxide below 100 microns, you can go to practically 19 anywhere it's willing to support. 20 CHAIRMAN POWERS: As long as there's no 21 change in the physics, which is not demonstrated here. 22 Well, that's the next MR. MONTGOMERY: 23 I'm trying to demonstrate that through the slide. 24 the again, have, We experimental database. 25

experimental database for the conditions for which 1 this curve's applicable, which is 300 C, 280 C or 2 above, we only have these data points that have not 3 failed -- or that are not spalled, okay? 4 these had spalled oxide. They had oxides up to 100 5 micron but they did not have spalling. 6 We have tests out to 63, 64,000 that are 7 very near our curve and did not fail. We have this 8 one that's above our curve that did not fail. 9 then we have this one that's well above our curve, and 10 this one is a bit of -- I don't want to call it 11 anomaly, but in a sodium reactor you're not going to 12 post-DNB heat transfer conditions, so you don't really 13 -- can't really say that that's -- that the failure 14 could be moved that high, it's just that PCMI is not 15 active at that level of enthalpy to cause failure. 16 So here in this curve MEMBER ROSEN: 17 you've actually -- you've drawn out to 90 gigawatt 18 days per ton. 19 MR. MONTGOMERY: Well, yes. Just note --20 now, be careful here. 21 (Chatter.) 22 MR. MONTGOMERY: Be careful. Let me point 23 out something. I was obscuring it in my standing of 24 the -- where I was standing. Since heat is short

1	segments over 50 centimeters or so, they represent a
2	peak burnup per se, so this curve has been moved from
3	a rod average burnup to a peak burnup, and that's
4	where there's a shift to a higher burnup. So this
5	would be the largest the peak burnup, the peak
6	nodal burnup. For a rod at 75 average would be about
7	85, 86, 87 type number. Eighty-eight is what's
8	plotted here. That depends on the peaking factors of
9	the plant, the axial power shape. So that's the
10	difference between the two curves. This one is on rod
11	average basis, and this one is on rod peak basis.
12	MEMBER FORD: Just to make sure, Rep-5, 11
13	and 4 are no failure?
14	MR. MONTGOMERY: All of these have no
15	failure. And I should point out, just for
16	clarification, is that RepNa-11 is a UO2 rod but it
17	has the M5 cladding, it's got the more advanced
18	cladding, so it's oxide is really low, like 30
19	microns.
20	MEMBER FORD: But in Rep-2 not failed?
21	MR. MONTGOMERY: It did not fail because
22	at this low of burnup the oxide is rather low and the
23	ductility cladding is sufficient to accommodate the
24	loading from the pellet. It was tested in sodium so
25	you don't get the high temperature mechanisms of

oxidation-induced embrittlement that would occur if 1 you were to test this same type of test in water. 2 that's why it did not fail. Okay. 3 All right. So this is what I have to say 4 about the failure threshold criterion that has been 5 I will now -- unless there's some established. 6 questions about this, I will move into the coolability 7 discussion and talk about core coolability. 8 Just to make sure Ι MEMBER FORD: 9 understand, if you had oxide scoring, then at around 10 50 what you'd see is that you'd have a 11 about discontinuous curve and it should just drop down to a 12 13 value. MR. MONTGOMERY: Yes. The spalling curve 14 would be down here. 15 Down here, and it would MEMBER FORD: 16 presumably loop up to join that main curve. 17 loop up here. MONTGOMERY: Yes, MR. 18 Because when the spallation process kicks in, it's a 19 fairly -- there's a step almost change between the 20 ductility between spalled and non-spalled. 21 MEMBER ROSEN: So one way to supercondense 22 this discussion for us laymen is to say the transition 23 to advanced cladding materials is done to make sure 24 that you don't get thick oxide layers, so that you 25

1	don't have a potential for spalling, so that you don't
2	get hydride mobility which can lead to low ductility.
3	MS. SIEBER: And that protects you against
4	prompt pulses which you'll never get.
5	MEMBER ROSEN: That's right. All of that
6	work is to protect you against something you'll never
7	get. But if you did, if you could imagine it, you
8	would be okay anyway.
9	MS. SIEBER: You could do it but you've
10	got to put a tunnel in there to get it in there.
11	MEMBER ROSEN: All you got to do is just
12	ten percent more and you get the 100 megawatt days per
13	ton, which is where
14	MS. SIEBER: Just bigger paper. Once you
15	draw beyond the data, it becomes a matter of how
16	embarrassed you are.
17	(Laughter.)
18	MEMBER ROSEN: And for those of us who are
19	never embarrassed about anything?
20	MR. MONTGOMERY: Now we have a couple of
21	pieces of data that are going to come in in this range
22	right here, right, Rosa, for this step 1 and step 2
23	test. On M5 cladding, they'll come in on this range
24	in the next coming months.
25	Up to now we've been talking about failure

and the threshold required to define when it's necessary to start counting for radiological releases to meet the dose requirements. So the next subject that I'll move to now is the coolability concern, which really represents the safety limit with regards to maintaining the core geometry.

The database is a bit smaller in that regard than for the failure database. The past experiments in the U.S. and Japan early on focused on enthalpy generally above 280 calories per gram. primary objective was to look at molten fuel, kinetics the mechanical dispersal and energy generation from fuel coolant interactions. They really wanted to see what was happening at very high energies to understand the real safety consequences.

Recent experiments we've had in France and Japan generally have been below the 220 calorie per gram limit. You saw one point that I had from CABRI that was about 215, and we have a couple from NSRR that are on the order of 210 or so. And some of these cases and for those that experience failure, some of them have dispersed a small amount of finely fragmented solid materials generally coming from the pellet periphery.

And in some of these cases, there is a

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

measurable amount of mechanical energy generation, and 1 maybe I should just talk briefly about what I mean 2 3 about that. Particularly in Japan, they use a 4 stagnant water system where the fuel segment, again a 5 six-inch segment, sits in a canister with a -- in a And at the top of the water they 6 pool of water. generally put a float device, and the float device has 7 a magnetic sensor system on it so when the float bumps 8 up and down they can measure the velocity and how far 9 that float moves up and down. And what we mean then 10 11 by mechanical energy generation is that in the process 12 of conducting a test if they measure that float moving with some significant velocity and have some upper 13 movement and the height that it moved to, they can 14 then determine from the energy, mechanical energy 15 16 generation from that process. So that's what I mean 17 by mechanical energy generation.

The fuel dispersal is an issue. It occurs generally at burnups greater than 40,000 due to the rim effect. The increase in local burnup and fission density in the outer rim influences the temperature and the local effects that go on in this area and when the cladding fails can promote some material to be dispersed from the fuel rod through the cladding out to the coolant.

18

19

20

21

22

23

24

1

2

3

4

5 6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

Some of the issues that are raised as a consequence of fuel dispersal is if you can significant amount of material out, could it result in loss of low blockage or loss of raw geometry such that These are geometrical you can't maintain cooling? And then more of pressure vessel type effects. integrity issue is that you could get pressure pulse generation that could effect either the core geometry, again from a cooling point of view, or the vessel So this is something we need to integrity itself. look into. And so we've looked at the data and what we see is that the potential for fuel dispersal is a function of how much energy is deposited after the cladding has failed and also the pulse width.

So we've come up with a slide here that shows the data on high energy tests that have had cladding failure and post-failure energy deposition. So we have along the Y-axis here is the energy deposition after the cladding has failed, and plotted along the X-axis here is the pulse width. And you see that for most of these tests that were tested below ten milliseconds there is some fuel dispersal that occurs, and that's separated by the points on this side of the dash line all had some sort of fuel material -- solid fuel material dispersal, a few grams, generally, or less.

And then the tests on this side of the line, although they failed, had developed a crack in the cladding, none of the fuel was released from the -- cut from cladding and into the coolant. So that there is some -- you can see that there's some effect of pulse width and then effect of energy deposition after failure.

This very busy schematic illustrates the processes that are controlled by pulse width that can influence the dispersal process. Here in this illustration is the narrow pulse and in a narrow pulse, as I showed earlier, we get these temperature distributions where the peak temperature occurs in the outer pellet periphery region. As a consequence of the rapid energy deposition rate, the heat transfer conditions are slower so you don't have as much heat transfer. So you generally end up with higher temperatures in that region and stepper gradients in that region.

Combine that with the fission gas distribution and content and the pellet in that region, you can end up then with higher gas pressures and higher thermal stresses as a consequence of these gradients, and end up with the fuel tending to

fragment apart and what we call grain boundary 1 2 decohesion, resulting in fission gas release and also now that the fuel is fragmented a bit, it has the 3 4 potential to be dispersed. And this micrograph is a micrograph from 5 It's a sermography here. This is the fuel 6 7 pellet, this would be towards the center of pellet, and this is the pellet periphery. The 8 cladding would be just over here. It's kind of hard 9 10 to see, but there's kind of a gap right here. individual 11 what we see is that you see boundaries that are decorated, and you see there's a 12 crack here, there's a number of cracks here and here 13 And you can see that the grain boundaries are 14 very evident, and that indicates that the grain 15 boundaries have more than likely separated off and the 16 fuel is almost cracked up into approximately grain-17 size segments or bigger, on the order of ten to 20 18 microns. 19 There's a marked difference MS. SIEBER: 20 in density in the right third of that micrograph. 21 MR. MONTGOMERY: Here versus over here? 22 MS. SIEBER: Yes. 23 Yes. This is the rim MR. MONTGOMERY: 24

region.

It looks like a straight MS. SIEBER: 1 Could you tell me what that is? 2 line. MR. MONTGOMERY: Right here? 3 No, over to the left. MS. SIEBER: 4 MR. MONTGOMERY: Right here? 5 Right there. MS. SIEBER: 6 MR. MONTGOMERY: That's an artifact of the 7 etching more than likely. This is the rim region 8 where the grain size has decreased some, and when the 9 etching is done to generate this micrograph usually 10 that region where the rim is comes out stronger, 11 showing a stronger etching. So this is where the rim 12 generally is, and you get a finer grain density in 13 that area. And it's a pretty sharp transition between 14 It could be a photograph artifact as well. 15 the two. On the other hand, if we have a wider 16 pulse, generally on the order of 20 milliseconds or 17 greater, there's time for a heat transfer so the 18 temperatures tend not to go quite as high, 19 20 gradients are smaller. These combine together to have less of effect on the local gas pressure and the 21 bubbles and on the grain boundaries, and limits the 22 cracking fragmentation and the possibility of fuel 23 dispersal. And you can see here that these -- again, 24 this is RepNa-4, had a wider pulse, and we don't see 25

- 11	156
1	quite the level of cracking and grain boundary
2	separation in this micrograph. Again, this is the
3	pellet surface region and that's going towards the
4	center. We don't see quite the level of grain
5	fragmentation.
6	MEMBER ROSEN: And, again, that's the
7	artifact that Jack was talking about in RepNa-4, the
8	photograph.
9	. MR. MONTGOMERY: I'm trying to see.
10	MEMBER ROSEN: One side's very light and
11	one side's very dark.
12	MR. MONTGOMERY: These could be two
13	photographs here. Yes, that's a montage. Although
14	this is really not a montage here. It looks much
15	better on my screen.
16	So back to this prototypical pulse width,
17	for prototypical pulse widths no fuel dispersal is
18	expected. However, at high energy after failure, it's
19	possible that a small amount of non-molten pellet
20	material may be dispersed, but it's impact is low. We
21	have experimental data to support that. In a test at
22	NSRR approximately ten percent of the pellet was
23	released, and in that case the fuel rod maintained the
24	geometry.
25	I have a little slide I added that I

wanted to show you just to give a feel for what I'm talking about. I don't have it in a handout, I'll be happy to give it to you if you want it. On the top here is a rod that was tested up over 200 calories per gram, it developed an axial crack, you can see here, and then in further post-test examinations you can see the crack in the cladding. Here's the fuel pellet. It was pre-irradiated to about 30-something thousand, 30, 35,000, and you can see that there's some material lost right in the vicinity of where the crack is that some material has been released out. You can see a little bit of loss in this region here consequence of the test. And this test lost about ten percent of the fuel material was -- left the cladding and was found in the coolant. But the rod still looks like a fuel rod, and it's still maintaining a geometry that is coolable and contains a majority of the fuel material.

This is just a picture, I know you can't see this very well, but that's just a picture of the material that was found outside the fuel rod. You see small pieces. The key point here is that none of it looks molten. This test was done almost to the melting temperature, and it clearly did not reach that, and the material that left the fuel rod was not

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

molten. The difference would be is if this was molten
material, it would look like a bunch of BBs, pellets,
you know, round balls almost.

Okay. Generally, what we see is that for any fuel coolant interaction that results in pressure pulses is that the tests exhibit rather low mechanical energy conversion primarily because the material temperature is low and molten material, so it has less stored energy. And the heat transfer kinetics aren't as energetic. Secondly is that there's a limited amount of material, generally just a small amount of rim material is what's been released.

So to establish the coolability limit we elected to use an enthalpy limit that would preclude incipient melting so that if in the off chance some material is dispersed it would not be molten. As I said, the data show that dispersal molten material generally produces higher thermal-to-mechanical energy conversion ratios. I'll show that in the next slide. The test that I just showed you, this JMH-5 which was tested up at 200 calories per gram, showed no adverse impact on rod geometry. Even though it dispersed a small amount of material, it maintained a rod-like geometry. And that there would be no impact on the pressure vessel because the pressure pulse -- the

1 || mechan

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

mechanical energy generation would be low.

incipient melting fuel Using coolability limit is for the precursor conservative in the sense that we really are limiting below the melting fuel to well the of most If we define the peak temperature here temperature. to be below the melting temperature, a majority of the fuel is well below that because of the peaking effects It also limits such that the at high burnup fuel. cladding does not reach melting, so we maintain rod geometry in that fashion as well. And, finally, it limits the thermal-to-mechanical energy conversion.

And that's shown here where we're looking at -- this is a subset of the data done from the early Japanese and CDC SPERT tests where they tested the fuel up in the molten area. So they're all tests done about 320 calories per gram or higher. And you can here mechanical energy plotting that I¹m see conversation ratio versus the particle size when they look at the particles after the test. And we see that the data shows a dependency on the particle size and can get up to one percent energy conversion when the material is molten.

If we go to non-molten material tests, that's these two, and this is the real important test

because these are powder tests, these are done with powder, special tests done to simulate powder being dispersed, we see that the conversion ratio is about an order of magnitude less and the total energies that are generated are even larger than that between the two, if you look at the energy generated in this versus the energy generated in that. So there's quite a bit of difference between non-molten and molten material. And the dependency on particle size is much less. This generally has about a square root dependency on particle size, and this has about a linear dependence on particle size.

So in order to establish a limit on the enthalpy to preclude incipient melting, we need to determine what enthalpy would be necessary to reach the melting temperature. So to do that, we did an analysis again where we combine data on the UO2 melting temperature as a function of burnup, combine that with the radial burnup and power distribution information that we know to give us the local burnup and the local temperature, melting temperature. This gives us the local burnup and the local melting temperature, and then through an analysis, using a pulse width of 20 milliseconds, we calculated what the enthalpy would need to be to reach the melting

temperature locally in the fuel and then define that as the maximum enthalpy level as a function of burnup.

And we did this through the burnup range.

And the answer is shown here where we again, maximum radial average fuel enthalpy versus rod average burnup. This is the result of the A limit was placed on -- this curve analysis. actually goes up to about 250 or so, but I went ahead and just capped it at 230 because that's kind of where the licensing base of today is anyway. And what we see is I plotted out some of the data here from the zero burnup tests that have been done where we've had some maintained rod geometry here, we have clad melting in this range, partial clad melting, and then total loss of rod geometry, as indicated by these symbols. And then I've overlaid the few tests that are in this energy range of interest where they've been tested up to about 200 calories per gram or so. That's where this database is. And all those maintained rod geometry. So that's the data compared to the limit line.

So I'm getting close to being finished up here. You saw this curve before, Rosa showed it. What we have here is the failure threshold as a function of burnup and the core coolability limit as

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

a function of burnup. They incorporate the effects of burnup through the material properties and melting temperature.

To summarize, we've proposed acceptance criteria, fuel rod failure threshold and the core coolability limit, that as a function of burnup we think that these acceptance criteria include the key controlling parameters, that is the corrosion and hydriding evolution with burnup that affects the material ductility and failure and the burnup impact on UO2 melting. These criteria are given in terms of radially averaged peak fuel enthalpy. This consistent with the current reload design methods where the neutronics calculations generally calculate this parameter as one of their outputs. Currently, it's applicable to hot-zero power RIA events. At this point in time, we feel that DNB remains the limit, the appropriate failure criterion for at-power rodejection events.

The failure threshold is based on integral tests from RIA simulations, mechanical property tests and analytical methods. It's certainly based on the corrosion kinetics that we used. It certainly represents a lower bound for modern, low corrosion cladding. And, as I said, it tends to bound the data

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

for the non-spalled Zirc-4 rods.

For coolability limit, we don't expect any fuel dispersal to occur during LWR conditions, but if there is, we've put a limit on the peak fuel temperature or the enthalpy -- put a limit on the enthalpy to preclude incipient fuel melting. This is now a function of burnup. And it is supported by data from the database that we have on both loss of rod geometry, mechanical energy release. We feel this is conservative and in general we get much less than ten percent of the fuel material that's going to come out. And then there's a large margin between peak burnup that we assume in this calculation and generally the location of the peak energy deposition, and that's given in the next slide.

burnup distribution from a high burnup rod, you can see that the burnup is about 55,000. And superimposed on that this is burnup on this axis versus axial position. And we have superimposed on that the axial power shape during a rod ejection event, and in a rod ejection event the axial power shape is very much peaked in the top of the core because of the characteristics of the event. And we see that for this case the axial peak power, which we assume in our

analysis, this would be the radial average peak. 1 2 (Whereupon, the proceedings went off the 3 record at 12:31 p.m. and resumed at 1:32 p.m.) 4 CHAIRMAN POWERS: Let's come back into 5 session. Undine, you are going to tell us about what you are going to do about all this good stuff we have 6 7 heard about, right? MS. SHOOP: Absolutely. You are going to 8 9 be dazzled and impressed. Okay. I would like for the 10 second part of this presentation --CHAIRMAN POWERS: We are always dazzling 11 12 and impressive. I won't go there. I would 13 MS. SHOOP: like to talk to you how we are actually proposing what 14 kind of plan we have come up with, a preliminary plan 15 16 actually, to review this topical report. 17 The purpose of generating a plan to begin with focus to 18 is that we can our resources appropriately provide detailed review, 19 the and 20 identify all the elements up front so that we make 21 sure that we are not missing anything, and that we complete review and that there 22 are no 23 surprises. This is a team effort. There is myself, 24 Shi-Lang Wu, and Ed Kendrick on the NRR team; and then 25

we are also working with the Office of Research, and 1 also Carl Beyer from PNNL, our contractor, provides 2 support for this. 3 The elements of the review plan currently 4 include data verification. As you have seen, there is 5 a lot of databases, and there is a lot of databases 6 from a lot of different tests. 7 And what we need to do is make sure that 8 all of the data is applied in a manner consistent with 9 the way that it was generated. It is applied and 10 there is a correct application of the methodology, and 11 any time that you get more than one task, you always 12 have uncertainties. 13 So we need to make sure that all of the 14 data is in line. 15 CHAIRMAN POWERS: Do you mean to tell me 16 that with one test that we have no uncertainty? 17 MS. SHOOP: You can talk to Rosa about 18 19 that. MEMBER ROSEN: With two points, you have 20 a straight line, and with one point, you have the 21 answer. 22 I never CHAIRMAN POWERS: You know, 23 thought of that. You may have established a new 24 principle of science there. Don't expect Stockholm to 25

call too soon though. 1 So you know that Okay. MS. SHOOP: 2 statistics is always our favorite thing, and so we are 3 going to look at that as well. In the SED/CSED theory 4 and model, we need to investigate, and come to terms, 5 and verify ourselves that the SED/CSED model is an 6 equivalent of Rice's J/Jc formulation, which was the 7 inter-role of the strain. 8 And then we are going to code the SED/CSED 9 formulation into the NRC FRAPTRAN code. That way we 10 can do an independent verification of the analysis 11 results that EPRI has presented. 12 In the fuel rod failure thresholds, we are 13 going to have to validate the application, and we are 14 also going to have to review it for applicability to 15 the current future and proposed fuel types just to 16 make sure that everything is bounding. 17 In the core pool ability limit, we have to 18 do application verification. As we have seen today, 19 there is some limited data, and then some of it is 20 from analytical methods. 21 And we need to make sure that that is 22 rigorously addressed and appropriate. The FALCON 23 code. EPRI uses the FALCON code in the development of 24

this methodology, and that is a code that the staff

1	has not seen, nor have we ever reviewed it.
2	And EPRI has graciously agreed to provide
3	us with a copy of the code. That way we can look at
4	it and review it. The data dispersal
5	CHAIRMAN POWERS: Will they be giving you
6	things like users manuals, and models and
7	correlations, and things like that?
8	MS. SHOOP: Yes. They are exceptional
9	gracious. They are providing us training with the
10	code, and they are providing all of the V&D, users
11	manual, and the theory manual, as well as the source
12	code.
13	CHAIRMAN POWERS: Are you going to share
14	it with us?
15	MS. SHOOP: What are you guys going to run
16	it on?
17	CHAIRMAN POWERS: What do you mean? I
18	have access to a computer with 3,000 processors, 1
19	gigabit, 1 gigahertz each node. Is that enough?
20	MEMBER ROSEN: That ought to be enough.
21	MS. SHOOP: I think I am going to log into
22	that machine. In the area of field dispersal, where
23	you have to review the data for applicable at each of
24	the phenomena of the proposed safety limits, there
25	again do the validation and verification.

For the uncertainty and conservatism, you 1 know, we always have to look at the uncertainty, and 2 3 we have to verify that the conservatism is appropriate But for the limitations of the and bounding. 4 criteria, you have to review the data for where it is 5 applicable, and make sure it is applicable for the 6 full range that we anticipate it being used for. 7 safety And then also have our 8 we evaluation conditions of acceptability. 9 have those. And what type of fuels are applicable to, 10 and is there any sort of core design limitations, or 11 anything like that. We will of course always look 12 into that. 13 This is also going to entail revising the 14 Reg Guide, Reg Guide 1.77, and also there is three 15 SRPs that all reference this limit. And they will all 16 have to be verified. 17 So of course we will be coming back to see 18 your smiling faces to show you the reg guides and the 19 SRPs, and get your weigh in after all of this is all 20 Okay. 21 done. CHAIRMAN POWERS: This is the highlight of 22 your schedule, right? 23 MS. SHOOP: Yes. 24 CHAIRMAN POWERS: That's good. 25

Coming down to see you MS. SHOOP: Yes. 1 Since this is a team guys is always a highlight. 2 effort, we will as I alluded to before, we will be 3 asking the Office of Research for some assistance. 4 The Office of Research is very familiar 5 with the data, and with the testing mechanisms, and so 6 we will definitely need their assistance with 7 verifying that the application methodology is applied, 8 and all data is used consistently. 9 The Office of Research also has a contract 10 for the FRAPTRAN computer code. So incorporating the 11 CSED/SED model into the FRAPTRAN code will entail 12 getting their assistance in that respect. 13 And I should actually back up, because 14 That will be a DPRI is looking a little worried. 15 proprietary version of the FRAPTRAN code and that will 16 not be a publicly available one. 17 Fuel dispersal. We are going to also ask 18 for their assistance with the applicability of the 19 fuel dispersal the proposal for the 20 to mechanisms. And my last slide. 21 Our offices, since this is a preliminary 22 review plan, we are planning on coming up with an 23 office agreed upon final review plan, and we 24 anticipate having that by December of this year. 25

1	you have any questions on our review elements or our
2	proposal? Rosa has a question.
3	MS. YANG: May I ask just for
4	clarification, because we were given a schedule of the
5	review, and does this bullet mean that you may revise
6	that schedule?
7	MS. SHOOP: The schedule may actually
8	you know, there were some interim dates in that
9	schedule, and they may move as you know, some of
10	those we discussed with the Office of Research, and
11	then some of them we have since gotten input.
12	So we need to dialogue between our offices
13	and see if any of those interim bullets need to move.
14	MEMBER FORD: This is saying that you must
15	have already done many of these tasks.
16	MS. SHOOP: No, this is just saying that
17	we are going to come up with a final plan on how we
18	are going to review this topical by December.
19	MEMBER ROSEN: No. That's not the finish
20	date.
21	MS. SHOOP: We have started to review, but
22	that basically will lay out the elements of the
23	review.
24	MEMBER FORD: That would be wonderful if
25	we could have it by the end of the year.

MS. SHOOP: And it will be in a laid out 1 plan that both offices agree to. 2 So when will the final MEMBER FORD: 3 review be done? 4 CHAIRMAN POWERS: Whenever the -- it says 5 on December 31st that they will answer that question. 6 Okay. Well, thank you, Undine. 7 I am putting my mouth in MEMBER FORD: 8 EPRI's foot, or my foot in EPRI's mouth, but I would 9 imagine that they would want this to be done fairly 10 quickly. Is there any way of pushing it up, or is it 11 not high on the prioritization, or what? 12 of There are a number MS. SHOOP: 13 components, and yes, and any licensee who comes in 14 here with a licensing application wants it done 15 quickly. That is just a given. With this particular 16 plan, this is one of a series of different topicals 17 that will have to be submitted to support high burn-18 up. 19 And high burn-up is of interest to the 20 and it is important to the agency, 21 therefore we need to make sure that we take the 22 appropriate time and resources to do a thorough review 23 to have all of our ducks in a row to approve it. 24 MEMBER FORD: This also means that you 25

1	will be doing therefore on the high burn up fuel,
2	which is contrary to the message that we were getting
3	before.
4	MS. SHOOP: We will be doing work in the
5	areas of reviewing what the industry has provided.
6	MEMBER FORD: Yes.
7	MS. SHOOP: Keeping abreast ourselves of
8	what is going on in the international community, and
9	to see how that all relates. However, we are not as
10	I said in the Agency's 1998 plan, we have said that
11	the onus of coming up with the criteria database and
12	the methodology will be the industry's responsibility.
13	MEMBER ROSEN: And they have done it.
14	. MS. SHOOP: Yes.
15	CHAIRMAN POWERS: Okay. Thank you. Well,
16	now we are going to switch gears a little bit and move
17	to the question of the RES program. And I guess we
18	are going to start with Jack, who is going to give us
19	
20	MR. ROSENTHAL: My name is Jack Rosenthal
21	for the record again. I just wanted to say that this
22	is a very good time when we welcome coming here, and
23	we are trying to generate test data relative to LOCA,
24	and Argonne.
25	We have finished some Surry creep data

that is important in the waste arena, again with some 1 data, and so after many years of promising, we are 2 finally seeing some results. So it is a very good 3 time to brief you on where we stand. 4 And Ralph Meyer will go over what the 5 promises were from 1998, in terms of the plan, and 6 what we have accomplished, and where we are. And then 7 you will hear more detailed presentations mostly on 8 our experimental program. 9 CHAIRMAN POWERS: Ralph, I'm sure that you 10 are going to say this, because I have looked at your 11 want to reiterate that to this Ι slides, but 12 subcommittee that we have looked in great depth at the 13 reactivity insertion accident aspect of high burn-up 14 fuel. 15 There are many other aspects of high burn-16 up fuel impacting issues of safety, and I am sure that 17 Ralph will touch upon at least some fraction of those. 18 Ralph. 19 MR. MEYER: Yes. Actually, we have four 20 hours of presentation prepared, and we will shorten it 21 up, and quit before the sun goes down or something. 22 Anyway --23 CHAIRMAN POWERS: Who imposed this sundown 24 criterion? This Committee is used to being here until 25

7:30 or 8:00 o'clock at night. 1 MEMBER ROSEN: It is normal ACRS practice 2 to say when someone who says they have four hours to 3 give them 40 minutes. 4 CHAIRMAN POWERS: Go ahead, Ralph. 5 MR. MEYER: Okay. We are in fact working 6 on a revised, or you could think of it as a new high 7 burn-up program plan that would cut across the 8 offices. 9 CHAIRMAN POWERS: So we have heard. 10 MR. MEYER: In addition to the plan for a 11 review of a particular licensing topical report, there 12 is a broader update in progress, but we are not 13 finished with that. 14 So what I thought I would do would be to 15 roll back to the 1998 plan, and tell you where we are 16 on the issues that were identified in that plan, 17 because the new plan will obviously pick up and go 18 forward in some manner on these or other issues. 19 So here is the original list of issues, 20 and just to identify, there were nine of them; 21 cladding integrity, control rod insertion problems, 22 reactivity accidents, which we have talked about all 23 loss of coolant accidents, the morning; 24 oscillations in BWR associated with an anticipated 25

transient without SCAM. 1 Our computer codes for fuel rod behavior, 2 and neutron kinetics; a source term for high burn up 3 fuel, transportation and dry storage issues related to 4 high burn-up fuel, and high enrichments. 5 Now, in 1998, we said that Issues 1, 2, 6 and 9 were essentially either resolved or we didn't 7 need to talk about, and so I am not going to talk 8 about them today. I am going to concentrate 3 through 9 10 MEMBER ROSEN: It is a good thing you have 11 four hours, because let's talk about some of those. 12 And I want to talk about nine in the context of the 13 advanced reactor research plan that we are working on 14 here. 15 If we are really serious about -- if the 16 agency is serious about writing research, an advance 17 reactor research plan that considers the introduction 18 of fast reactors, either gas cooled or liquid metal 19 cooled, or in any, you are going to need enrichments 20 greater than five percent. 21 MR. MEYER: Yes. 22 So there is some sort of MEMBER ROSEN: 23 something going on here and I don't know what the --24 I am just rolling out the rope here. 25

1 MR. MEYER: We have work in place to look at advanced reactor fuels. We have an advanced 2 3 reactor research plan that has been developed that includes both fuels, and therefore, would include 4 5 higher enrichments. But in the context of high burn-up fuel, 6 7 the industry has decided that it would like to make additional steps in increasing burn-up, but that they 8 would not need to go beyond 5 percent enrichment in 9 10 current light water reactors in order to do that. So in terms of a program plan that is 11 12 looking at high burn-up fuel in current reactors, it is pretty much off the table for us. 13 did provide 14 MR. MONTGOMERY: We 15 advanced reactor research plan. You know a draft plan to the ACRS, like I think two days ago. So you ought 16 to find it appearing in the in-boxes shortly. 17 And the big thing was to add ESBWR and ACR 18 19 700 to that plan, and they are not high enrichment. The ESBWR, for example, uses modern boiler fuel. IRIS 20 is out in the distant future, and at one time we 21 thought that that would involve high enrichment fuel 22 23 when they were talking about multi-year cycles. οf Thursday, last Thursday, at a 24

presentation that they made here at the NRC, they

1	indicated that at least for now that they did not plan
2	to go above the five percent enrichment value. So
3	that is where we stand right now.
4	MEMBER ROSEN: Well, we will duly note
5	that, and the advanced reactor research plan comments
6	that this committee will offer some time. It is
7	pretty clear that you can't get there from here for
8	that comprehensive list of things that are apt to be
9	in the plan, or apt to be on the table.
10	At least they are in the Gen IV list, and
11	in the international and near-term deployment list.
12	They may not be in the domestic near-term deployment
13	though. There are enough concepts in those lists that
14	will require enrichments beyond five percent and that
15	somebody in the agency ought to be thinking about,
16	rather than just dismissing it out of hand.
17	I understand that in this case that you
18	are dismissing it out of hand because this is a plan
19	for the current light water reactors.
20	MR. MEYER: Yes.
21	MEMBER ROSEN: And I agree that nobody is
22	talking about greater than five there.
23	MR. MEYER: Shall I go on?
24	MEMBER ROSEN: Yes.
25	CHAIRMAN POWERS: Please.

Okay. Now I want to talk 1 MR. MEYER: about several of the issues, including the reactivity 2 I plan to do this by way of an 3 initiated accidents. introductory presentation and then a revisiting of the 4 5 issues in the subsequent presentations, and to go into a little more detail about work that we have actually 6 7 done. So it is somewhat of an artificial split, 8 9 and there is likely to be some interest in jumping into the second presentations right now, and we can do 10 11 that if you want to, or not do that. CHAIRMAN POWERS: The subcommittees are 12 controllable, but we will try and kind of constrain 13 ourselves and get a guick overview, and then delve 14 into the details. 15 MR. MEYER: Okay. 16 CHAIRMAN POWERS: So just feel free to say 17 stop, and I'll tell you about this later. 18 Well, the issue was MR. MEYER: Okay. 19 20 described well this morning, and it has to do with a regulatory guide number that we don't believe applies 21 to high burn-ups. I am going to show you in a diagram 22 on the next slide the method that we are going to use, 23 or the methods that we are going to use to address 24 this. 25

But before even doing that, I want to point out the schedule that we are working on. As Rosa mentioned, there are 2, or perhaps 3, CABRI tests in the sodium loop coming up in September. I'm sorry, in October, November, and perhaps a follow-up test early next year. It's not clear.

These are tests on ZIRLO and M5, and at the Argonne National Laboratory, we will be completing a series of mechanical properties test next year on high burn-up Zircaloy-4.

And there is a test in Japan that we are looking forward to, to try and get a handle on the temperature effects. You saw this morning that the Japanese tests were run at approximately 25 degrees centigrade, which is not the right temperature for the accident that we are thinking about.

And the Japanese have or are constructing a high temperature-high pressure capsule, which they expect to start testing in in 2004. And so our plan for providing a confirmatory assessment for Zircaloy clad fuel at 62 gigawatt days per ton is to wait for these tests, and give ourselves two years to get it all together, and in early 2005 come out with an assessment document that gives a story on why everything is okay with regard to reactivity accidents

for the current zircaloy fuel in operating reactors up 1 to the burn-ups that are licensed at this time. 2 CHAIRMAN POWERS: And this again is just 3 reactivity accidents here? 4 That's right. Ι have MR. MEYER: 5 different schedules for different things. But we are 6 now down to the point where we are talking about 7 fairly finite periods of time and definite schedules, 8 and definite activities. 9 CHAIRMAN POWERS: Now, there are a series 10 of CABRI tests scheduled to begin somewhat after or in 11 late 2005, I think? 12 Yes. MR. MEYER: 13 And so how do they CHAIRMAN POWERS: 14 figure in? Are they confirmatory of confirmatory? 15 MR. MEYER: Yes. In fact, that is the way 16 The program has been that we are looking at them. 17 delayed and they water loop tests themselves don't get 18 So I think we and underway until late 2005 or 2006. 19 EPRI have pretty much decided that we want to make our 20 decisions without waiting for that, and hope that 21 everything pans out according to those confirmatory 22 23 tests. It took too long to hold things up for 24 that, and I think we are learning enough that we can 25

go ahead and get much of the job done before then. 1 This is the same bunch of data that you saw before, 2 plotted in a different way. 3 4 This our so-called paint brush slide, and you won't be surprised to learn that we have a 5 view of the data and different 6 somewhat implications of the data than EPRI has. 7 So the picture that I am going to describe 8 is a little different than you heard this morning. 9 The first thing to notice is that we have plotted this 10 as a function of oxide thickness rather than as a 11 function of burn-up. 12 Obviously, the enthalph increase that a 13 fuel rod can withstand before the cladding breaks is 14 a function of several variables. You have talked 15 They re temperature, and rates which are about them. 16 related to pulse widths, oxidation, hydride, and burn-17 18 up. I think the oxide thickness has a stronger 19 effect than the burn-up has, because it directly 20 And so we have affects the cladding properties. 21 chosen to look at the data as a function of oxide 22 thickness, which does not have burn-up directly 23 associated with it. 24 So in that sense there is no limit out 25

here at the end in burn-up. This brings the scatter 1 down a little bit, but clearly doesn't remove the 2 3 scatter. Now, there are certain bodies of data in here whose 4 personalities we know a little bit about. 5 The Japanese data points probably should 6 be shifted upwards because the test temperatures were 7 too low. These CABRI data points should have probably 8 shifted downward because the pulse whip was too large. 9 And we will talk about this in the second 10 presentation in a little more detail. What I want to 11 say about this slide right now is that at the low 12 corrosion end of the plot, which is the low burn-up 13 end of things, the original correlation did indeed 14 have a relation to incipient melting. 15 The enthalpy for melting UO2 16 calories per gram, and if you do that on a radial 17 average, 230 calories per gram, is about where you 18 start melting fuel somewhere inside the rod. 19 There is a large volume increase going 20 from solid UO2 to liquid UO2, and this provides a 21 mechanism for breaking the cladding and expelling fuel 22 out of the rod which is what you saw or what we saw in 23 the earlier data at the higher enthalpy levels. 24 Now, 230 would be somewhere in here, and

you do see some cladding failures below that point, but you didn't get fuel dispersal in those cases because there was no mechanism for getting the fuel outside of the cladding. The cladding just broke open in some splits.

There is a big difference when you get to the higher corrosion rates which correspond to a higher burn-up, and there is definitely a correlation between burn-up and corrosion and Rosa showed one, or Rani did int heir presentation.

When you get the high burn-ups, and you heard this this morning, but I will just repeat it, you have this gassy grain structure in the fuel pellets. So now when you have a sudden temperature increase from the reactivity insertion, you have a rapid gas expansion, and you have a mechanism built in to disperse fuel if you can crack the cladding and produce some opening in the cladding.

CHAIRMAN POWERS: Let me ask you a question that might be better directed towards one of your subsequent speakers, and if so, I will be glad to wait. But when they do these tests, they cut out a section of irradiated fuel, and they put some fancy things on the end of it, and they may even repressurize it.

But when they cut it, they clearly lose 1 the gases that were in the nominal fuel clad gap, and 2 that was in the plenum. How much of the gas do they 3 lose out of this gassy structure at the perimeter of 4 the fuel that you are talking about? 5 MR. MEYER: Well, I don't think that they 6 lose any of that gas, because what we are talking 7 about is what we think of as non-released fission 8 gases, which are accumulated in tiny little bubbles 9 that attach themselves ot the grain boundaries. 10 And in high burn-up, you get so much of 11 that that it actually causes the grain boundaries to 12 subdivide a little bit. So you have got a relatively 13 fine grain material that has got a lot of these gas 14 bubbles on the grain boundary. 15 And I don't think you lose much or any of 16 that during the refabrication process. 17 Now that or those gas MEMBER ROSEN: 18 bubbles, micro bubbles, they don't form at the grain 19 boundaries exclusively do they? 20 MR. MEYER: The fission gases are not 21 soluble in the matrix and so they precipitate into 22 little bubbles, and the bubbles move around. And when 23 the bubbles get to a grain boundary, they share 24 surface area, and it is a lower energy position, and 25

1	so they stay there.
2	So what you predominantly see is all this
3	gas gets attached to the grain boundaries. So a large
4	inventory of the fission gases that have been
5	generated end up on the grain boundaries.
6	There is some still in the grains trying
7	to make their way out, but this is where they
8	accumulate.
9	DR. SIEBER: And some go to the plenum,
10	correct?
11	MR. MEYER: Some, not a lot, because the
12	well, 50 percent.
13	DR. SIEBER: So, 3 to 5 inches, and it
14	goes to a pressure increase of about a hundred pounds
15	over a 12 foot
16	MR. MEYER: Yes.
17	MEMBER ROSEN: Tell me again why does the
18	gas form within the grains, and migrate to the grain
19	boundary?
20	MR. MEYER: It is just a random process.
21	MEMBER ROSEN: It is a random process?
22	MR. MEYER: Yes.
23	CHAIRMAN POWERS: There is in fact
24	there is a thermal chemical driving force, two of
25	them. One is the temperature of the radiant, and the

1 other is --MEMBER ROSEN: It is a random process, and 2 the gas migrates around and when it gets to the grain 3 boundary, it stays there? 4 CHAIRMAN POWERS: It's not random. 5 MEMBER ROSEN: Any more. 6 MR. MEYER: When the first gas atom gets 7 in there, it moves randomly. It meets another one, 8 and they get together, and when you get a double, it 9 not a random process any longer because 10 This is an old temperature gradient gets involved. 11 story. 12 And we see them, and we believe they can 13 have this effect of pushing the fuel out through the 14 cracks in the cladding, because we have seen this kind 15 of dispersal in a number of the tests. 16 Now, this morning, you heard that typical 17 pulses in a PWR would be 30 milliseconds or bigger, 18 and they showed some data that showed that you only 19 saw this dispersal when the pulse widths were 15 20 milliseconds or less. Do you remember that slide? 21 22 Okay.

enough to fail the cladding will have a pulse width of

10 milliseconds or thereabouts.

Every PWR pulse that has an energy high

23

24

25

They will not be

1	broad.
2	The broad pulses that were spoken of this morning are
3	pulses with energies that are very low; 25 or 30
4	calories per gram or less.
5	If you get in the range where you can do
6	damage to the cladding, you already have narrow enough
7	pulses, except in a test reactor, where you can
8	contrive to make them broad, to expel fuel.
9	DR. SIEBER: Well, have you done any
10	research to decide what the pulse width will be in an
11	RIA in a reactor?
12	MR. MEYER: Yes. I will show you that in
13	the third presentation.
14	MEMBER ROSEN: And I thought you were
15	going to finish that sentence, Jack, in a real what?
16	DR. SIEBER: In a real reactor?
17	MR. MEYER: Yes. The answer is yes, and
18	I have a
19	DR. SIEBER: And is this a realistic
20	calculation or a licensing calculation?
21	MR. MEYER: That is a realistic
22	calculation.
23	DR. SIEBER: And you are going to show me
24	them?
25	MR. MEYER: I am going to show you them in

1	the third presentation.
2	DR. SIEBER: Okay.
3	MEMBER ROSEN: The third presentation? I
4	have to wait for that.
5	MR. MEYER: You have got to wait.
6	DR. SIEBER: Yes. Tell us when.
7	MR. MEYER: Okay.
8	MEMBER FORD: But the value of this plot,
9	paint brush thing there, is that your technical basis
10	for a specification of some sort?
11	MR. MEYER: No. This is just to guide
12	your eyes along roughly where the fuel failure level
13	is seen in these data. Now, for a PWR, we let me
14	get my story straight. Because of the potential for
15	a fuel dispersal here, we have chosen to take cladding
16	failure as the coolability limit.
17	In other words, whereas at low burn-up, we
18	recognize that cladding failure did not cause fuel
19	dispersal, and therefore, any consequences of fuel
20	dispersal.
21	So we worked with two different limits; a
22	coolability limit at a higher enthalpy, and a cladding
23	failure limit at a lower enthalphy for the purpose of
24	doing some dose calculations.
25	At high burn-up, we have chosen to

collapse this and to work on the cladding failure limit as the cladding failure threshold as the coolability limit. And I am confident that this is going to work because we are going to end up with cladding failure enthalpies in this range, somewhere in the 80 to 100 calorie per gram range, which is roughly twice as high as the enthalpy that you can deposit with a PWR experiencing a rod ejection accident.

And so it is a success path in my opinion, and I am going to show you how we are going to put it all together on the next slide. So we are searching for a curve that looks something like this, and what we are going to do with that is to do some plant calculations, which we can do, with a nice 3-D neutron kinetics code called PARCS.

And we are going to do some generic calculations looking at road works that are necessary to get you up to this cladding failure limit, and then by comparison with rod worths that are known from cordizines for commercial reactors to show that you don't have enough worth to fail the cladding.

And therefore none of the consequences that we are concerned about will take place, and that will be the end of our confirmatory demonstration.

1 || Now --

MEMBER FORD: There is a big assumption, and that is that your logic tree coming up with the results, and the big assumption is that the oxide thickness is the predominant metric of fuel cladding failure.

And if you can show that, great, but that is a dominant one, and burn-up has got nothing at all to do with it, and strain rate has got nothing to do with it.

MR. MEYER: Well, we are not -- I don't think we are constrained to saying that this is the only variable involved. Just as EPRI was not constrained to say that burn-up was the only variable involved when they plotted burn-up along this line.

In fact, we are going to be using correlations and codes that take many variances into effect in getting there. Maybe it won't look exactly like that two years from now, but this is the concept.

And we have actually three approaches, or maybe it is 2-1/2 approaches, to arriving at this -- at such a correlation. One of them is strictly empirical. You look at the data, and look at the various parameters, and try and correlate them.

Now, there is now for the past year a

correlation out there that we intend to work with. 1 Carlo Vitanza has developed a correlation based on the 2 CABRI data and the NSRR data, and I will show you a 3 little bit about that this afternoon. Not a lot. 4 So we intend to work with Carlo and 5 utilize a straight empirical approach from the data to 6 try and get some a correlation. We also have a fuel 7 rod behavior code, and can do in fact exactly the same 8 type of calculation that EPRI is doing with FALCON. 9 We can right now calculate the strain 10 energy density. It is just the integral of the stress 11 strain curve, and we can calculate stresses and 12 strains. Now, we are not as good at it as EPRI yet, 13 and we don't have a big an effort as EPRI or the 14 French have in the analytical area. 15 But we have got some improvements that we 16 are looking forward to soon, and that is one approach 17 that we can take. Another approach is just to look at 18 the individual data points themselves and move them 19 around on that plot based on things like temperature 20 variations. 21 I will talk a little bit about that. 22 think we can make some progress doing that. So we are 23 going to try 2 or 3 ways of coming up with an 24 enthalpy, a failure enthalpy curse, to be used in 25

confirmatory make our assessment, our to 1 order 2 assessment. Now, Ralph, I have been MEMBER ROSEN: 3 thinking about what you said, and it seems to me 4 aren't you getting a little ahead of yourself by 5 saying that it is a success path? 6 Because you could go through all of this, 7 assuming that you can do it effectively, and end up 8 with rod worth limitations that are so stringent that 9 nobody could design a cycle. 10 MR. MEYER: Well, you know, if we didn't 11 have a clue as to where we were going that would be 12 the case. But I can tell you right now that it looks 13 right now like the rod worths that you need to get to 14 cladding failure are about two dollars. 15 MEMBER ROSEN: Right. 16 CHAIRMAN POWERS: Jack, you had 17 question? 18 I'm trying to think Yes. DR. SIEBER: 19 about the practicality of the box on that drawing. 20 There is a correlation that EPRI put forward, and 21 perhaps it is an amorphous to some extent that equates 22 oxide layer thickness to burn-up. 23 And I am thinking as a plant operator 24 saying I really don't know what the oxide thickness is 25

of my core, nor do I really have the means to measure 1 it during a refueling. One thing I know is what the 2 3 burn-up is. So it seems to me that from a practical 4 standpoint that I would like to calculate burn-up 5 related to the oxide thickness, and then use that 6 correlation to determine whether I am in bounds or out 7 8 of bounds. And to me it is a more practical approach 9 and one which EPRI has chosen, too. 10 MR. MEYER: Well, we are not proposing 11 this for industry use. If you recall, we accepted an 12 obligation for the NRC to do confirmatory assessment 13 for current plants 62 gigawatt days per ton. 14 DR. SIEBER: Yes. 15 MR. MEYER: And what the industry does to 16 go from 62 to 75 is to be determined. I mean, there 17 is a proposal on the table, and a review under way. 18 this Yes, but with DR. SIEBER: 19 methodology, even at 62, you have got to make the 20 relationship between the oxide layer thickness, which 21 to me would vary from plant to plant, depending on how 22 the plant is operated, to the burn-up. 23 MR. MEYER: Well, we know that the oxide 24 thicknesses aren't much more than a hundred, and we 25

1	have got data out to like 130 in the database. And we
2	also have burn-ups in the database up to and a little
3	higher than 62. So I think we have covered the range
4	of the population of plants that we are trying to
5	address.
6	And so we will just go way out here to
7	where we think it is not any higher, and do the
8	calculation, and all indications are that we are going
9	to have ample margin to show that everything is okay.
10	CHAIRMAN POWERS: Yes, your objective is
11	to show that the decision to limit burn-ups to 62
12	gigawatt days, as opposed to 55 gigawatt days, still
13	provides adequate margin.
14	MR. MEYER: Yes. I would probably phase
15	it a little bit differently.
16	DR. SIEBER: Or 75, or 80.
17	MR. MEYER: But the approval is up to 62,
18	and for those approvals, we can demonstrate that we
19	have adequate margin for this accident.
20	DR. SIEBER: That's right, and in order to
21	approve that burn-up level though, somebody somewhere
22	has to make that correlation.
23	MR. MEYER: Well, it has already been
24	approved. This is after the fact. The 62 gigawatt a
25	day burn-up is approved.

1	DR. SIEBER: Well, what if you want to go
2	to 75? You still have to use the correlation to get
3	there.
4	MR. MEYER: Yes. Okay.
5	CHAIRMAN POWERS: Well, no, your
6	correlation is only good to 62.
7	MR. MEYER: Our correlation is as good as
8	their correlation. It is the same database, but we
9	are only attempting to apply it up to 62. I wouldn't
10	say it was no good above 62.
11	CHAIRMAN POWERS: You have got no
12	information.
13	MR. MEYER: What?
14	CHAIRMAN POWERS: You have got no
15	information right now above 62, or 65, or somewhere in
16	there.
17	MR. MEYER: Well, we are going to have a
18	couple of tests at 73 in a couple of months.
19	CHAIRMAN POWERS: But it is not your
20	obligation to defend a proposal to go to 75?
21	MR. MEYER: No, it is not.
22	CHAIRMAN POWERS: That's right.
23	DR. SIEBER: Well, I would suggest that
24	you go on while I ponder what you have said, and how
25	it fits into my working and thinking.

So now let me

Here we

1 MR. MEYER: 2 move on to the loss of coolant accident. wondered if the embrittlement criteria in 10 CFR 50.46 3 4 and the associated evaluation models either 5 Appendix K or whatever ones are being used, effected by burn-up, because in fact most of the 6 models and the criteria were based on data from low or 7

unirradiated materials.

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

Now, I have got several slides that I will show in a minute that talk particularly about the embrittlement criteria. So just to back up, and keep in mind that there were embrittlement criteria. 2200 degrees fahrenheit, peak cladding temperature limit, and 17 percent cladding oxidation limit. Those are the embrittlement criteria.

Okay.

Oops.

Then in Appendix K, or in a licensee's evaluation model, are some fuel related models. kinetics, and Oxidation a correlation for the occurrence of rod bursts, and a correlation for the amount of deformation in a ballooned section, and a correlation for a flow area reduction.

So these are the models and criteria that we are looking at to see what if any effect there is of burn-up. And we have work under way right now at Argonne National Laboratory, and harold Scott is going to talk about that in the second presentation.

We have Zircaloy-2 and Zircaloy-4 fuel rods with high burn-ups up at the laboratory, and the tests are under way, and it is our hope that in two years from now that we will have enough tests completed to be able to say something definitive about any changes in those criteria or evaluation models that we might have to make to accommodate the burn-up effects.

Now, I want to talk a little bit about the embrittlement criteria, the 17 percent oxidation limit and the 2200 degree fahrenheit cladding temperature limit. They arose as a pair of numbers, and they were related to some ring compression tests that were done by Hobson at Oak Ridge back in the late '60s and early '70s.

And the concept of a ring compression test was to take a piece of tubing, unirradiated tubing, cut some rings from that, and -- well, I'm sorry. First, oxidize a length of tubing.

And you oxidize it at some temperature for a period of time to accumulate a certain amount of oxidation on it. Now, what are the temperature ranges? During a LOCA transient, you heat up the fuel rod somewhere around 750 or 800 degrees centigrade.

1 The cladding balloons then burst, and then at a somewhat higher temperature, around 900 or 950 2 centigrade, the oxidation rate picks up, and as you go 3 4 up from -- let's say 900 to 1200, which is 2200 5 fahrenheit, and up to 1200 degrees centigrade, now you 6 are picking up a lot of oxidation rapidly. 7 So it is the amount of oxidation and the 8 temperature at which that oxidation is accumulated 9 that ends up becoming the embrittlement criteria. 10 in order to do that, you do a lot of ring compression 11 tests on specimens that have been oxidized 12 different temperatures in that range, and accumulated at different levels of oxidation in that range. 13 14 And you find that as log as the oxidation 15 temperature was not much above 1200 centigrade, and the total amount of oxidation is less than 17 percent 16 17 calculated by the Baker-Just correlation, then you 18 have ductility left int he ring specimen. 19 And if it is at a higher oxidation level, 20 you don't have ductility left, and this is the way 21 those two numbers were developed. So we are going to 22 try and replicate this process with high burn-up fuel. 23 But there is a little problem that came up

with all of this about a decade later, and that has to

do with some unexpected enhanced hydrogen absorption

24

on the inside of the cladding.

So here is a sketch of a rod that has ballooned and ruptured, and what was found was that steam got inside of the ballooned area and caused oxidation on the ID.

Well, that is taken into account in the regulation. We require ID oxidation to be calculated. So far, so good. The thing that wasn't so good was that the steam that was reacting with the zircaloy on the ID wasn't flowing. So the hydrogen that was released wasn't swept away.

And so a higher traction of the hydrogen that was generated on the ID got absorbed into the cladding, and now if you took a ring specimen from near the ballooned region, actually I am told that the effect is a maximum actually out of the balloon region and up in here.

And if you take a ring from that location, you find that at 17 percent, calculated by Baker-Just, it may be brittle when it was supposed to be ductile. Okay. There was some work done, and I just hasten to say that there was some additional work done at Argonne with pendulum impacters to show that in fact you still had ductility remaining for these specimens at the 17 percent level.

1 And so everything was under control, and this was all done back in 1980, 1989 or 1980, but we 2 3 have to understand this, and we are going to go back 4 through the process and do this kind of testing with 5 a high burn-up fuel. 6 MEMBER FORD: What is the justification 7 for doing the testing at 135 degrees centigrade? 8 MR. MEYER: What is the justification for 9 doing the testing? 10 MEMBER FORD: At 135. 11 MR. MEYER: Oh, oh, oh, yes. Now I am 12 going to forget exactly where the -- well, this is the 13 temperature at the end of the transient when you come 14 back down. 15 The Commission wanted ductility remaining. 16 There were big arguments about whether thermal shock 17 would fragment the fuel rods, and at the end of this 18 long hearing the Commission came down and said those 19 are all good arguments, but the only way to be sure 20 that we don't lose the geometry of the fuel rods is 21 that after this transient is all over, to have some 22 ductility left. 23 So this is the temperature at which it is 24 all over. These tests have been done at room

temperature as well, but I think 135 centigrade is

201 1 about where you expect the plant to be at the end of 2 a LOCA, a terminated LOCA accident. So that is where 3 they were done. 4 We probably are going to do them at 135 5 and at 20. But instead of doing pendulum impact tests to examine the effect of this enhanced hydrogen 6

> This is a fairly ambitious idea, but I think we will be able to do it. What this means is that we take a section of fuel -- of high burn-up fuel rods, with the fuel intact, and we sit it vertically in a channel, flow steam, and it is pressurized, and it heats up and we run it through a LOCA type transient.

> absorption, we are planning to do four-point bending

tests on ballooned segments.

And it balloons, and it ruptures, and it quenches, and it comes back down. Then we take the specimen out and we lay it down, and bend it in a four-point bend test, with the suspect region in the middle, and we let mother nature tell us where the weakest point is, and if there is any ductility left in this.

MEMBER FORD: The actual stress is in the component, and the stress would be highly biaxial, and highly anisotropic microstructure. And I am assuming

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

1	that someone has taken all of these into account, all
2	these aspects? You are applying a different stressing
3	mode to a
4	MR. MEYER: This is not I would like to
5	just say yes and hope that we went away from this, but
6	this is in fact about the right stress mode to apply
7	here, because you if have fuel rods experiencing
8	vibrations or seismic accelerations, they are going to
9	be lateral, and the fuel is going to be bending and
10	putting tensile stresses along the bowed out parts of
11	the parts of the road.
12	So I think it is just about right,
13	although I would say that we have a lot of work going
14	on looking at the biaxiality ratios, and
15	anisoptropies. These things tend to disappear at high
16	temperatures and at high burn-ups. So it is
17	definitely real critical.
18	DR. SIEBER: It seems to me when you get
19	down to about 130, or even as far down as 20, blowdown
20	loads are minimal.
21	MR. MEYER: Oh, blowdown loads aren't
22	really part of it, because the blowdown load is over
23	before the high temperature transient. What we are
24	thinking about are
25	DR. SIEBER: Seismic?

1 MR. MEYER: -seismic and just 2 vibrations, 3 and --4 DR. SIEBER: From what? 5 MR. MEYER: Well, unknown bumps in the 6 I mean, this is where the Commission in 7 debating this in 1972 and 1973, after discussing 8 possible loadings and talking about the actual 9 magnitudes of the loadings, decided that they couldn't 10 handle that analysis, and they said just give us some ductility when it is all over, and then we will be 11 12 happy. 13 MEMBER FORD: I think the argument goes, 14 Jack, that we will abuse this material as much as we 15 can, and then we will further stress it at the worst 16 possible temperature range, and see if anything 17 happens to it. It is a correlation to what could 18 really happen. 19 CHAIRMAN POWERS: That sounds like 20 something for PRA. 21 MR. MEYER: What we are trying to do is we 22 are trying to follow the spirit of the regulation as 23 it was originally defined, and simply investigate the effect of burn-up on that, without trying to reinvent 24

25

the whole procedure.

This is a sketch of the flow of the experimental work. You can think of another diagram 2 over here that is just a furnace with a piece end for 3 the oxidation kinetics measurements. 4 We have already completed a large series 5 of oxidation kinetics measurements, and Harold will 6 show you some of that later. It is just done in a 7 These are -- the furnace that we used is a 8 quadelliptical radiant heating furnace, heated from 9 the outside. 10 So ring compression specimens would be 11 12 13 14 15

1

16

17

18

19

20

21

22

23

24

25

prepared in a furnace, and we do the ring compression tests, and we look at the results, and we can do actual oxygen measurements, oxidation measurements afterwards, and look at the fracture surfaces to see if it is ductile or brittle.

Then in separate tests, we would do what we would refer to as a integral LOCA test, and from those specimens after doing a profilometry to look at the burst dimensions, then we can turn it over and do the ring compression tests.

And again look for the hydrogen, and do metallography on the fracture surfaces. So this is a general layout of the work that is going on. work has not started, but is expected to start very 1 | soon.

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

And we have done a substantial number of preliminary tests with unirradiated material, and in the last two months have completed two tests with high burn-up fuel rods.

And there is some rather interesting information coming, and just immediately coming from those tests which Harold will show you this afternoon.

The BWR power oscillations, this is not a design basis accident, and so what we are interested in here is whether or not a fuel rod that has gone into this situation and had the oscillations terminated in a way that the ATWS rule would specify that would consider this you successfully terminated.

And so in a successfully terminated ATWS with power oscillations, do you have benign behavior, or do you have non-benign behavior. I mean, in the PRAs up to now, it is always assumed that if you stop the oscillations in time that the behavior is benign.

But that was based on some analysis done by General Electric, where they used this 280 calorie per gram number, which is now not going to service well for high burn-up fuel. So it raises the question as to whether that assumption that we make in the PRAs

is correct.

CHAIRMAN POWERS: Well, even the process by which you get out of the oscillation is based on the assumption that there is ductility in the rods at sufficient level to withstand the dropping of the level, and remixing the boric acid when you have to, and things like that.

So, I mean, there is quite a lot that is involved here.

MR. MEYER: Let me try and -- oh, that one is in another one. I don't have any more on it in this presentation. I have a little more on this later. This work is going very slowly, and I tell you this every time. This is not our top priority, and we don't have a lot of horsepower working on this, but we have made some forward motion on it in the last year, and we will tell you just what little progress we have made in that presentation.

CHAIRMAN POWERS: Maybe this comes up later, but for us this is -- for the ACRS, and not for this committee or subcommittee, but for the ACRS as a whole, this is a great deal of interest to us because we are going through approving power uprates for boiling water reactors.

And when we come to the PRA and say what

is the risk significance of this, this is what -- the 1 issue that comes up, and the issue that comes up is 2 3 how shortened is the time available for the operator, 4 and if we go to high burn-up fuels as we well might 5 with uprated power reactors, are we getting into 6 regimes that are not. So I guess for us it is maybe a higher 7 priority than you see it in the fuel programs. 8 MR. MEYER: Okay. One of the next issues 9 10 on that list was our computer codes for fuel rod and neutron kinetics, and if you read the wording for the 11 issue, and what the issue was, it was related to the 12 fact that when we started reviewing applications for 13 high burn-up, our codes were not adequate for doing 14 audit work, because they had not been updated for high 15 16 burn-up analysis. And so we said as soon as we can get these 17 codes updated for high burn-up work, then this is not 18 an issue for us any more. It doesn't mean that we are 19 20 not going to do any work on our codes anymore, but it means that this particular issue would be resolved, 21 and it is now resolved. ' 22 The FRAPCON code was updated in 1997, 23 24 which is actually before the plan was issued, and the

FRAPTRAN report, the transient code, that was finished

25

(202) 234-4433

in -- well, just about a year ago, August-September of 2001 is when we issued documentation on the FRAPTRAN code with the high burn-up updates.

And in late 1998 the PARCS code was documented, and we are usually it routinely. So I think for the purpose of this plan that we have achieved our objectives for these code improvements, and this particular issue then ought to be considered resolved.

Source term for high burn-up fuel. I am almost afraid to say anything to this group about source term on high burn-up fuel. But the question was could we use the NUREG-1465 source term above 49 gigawatt days per ton, because in that report it said may not be applicable above.

There is sort of a bottom line on this, and the sort of bottom line is that we have met with a group of experts, and that elicitation was documented earlier this year, and if I could say it in a word, I would say that the 1465 source term is probably okay above 40 gigawatt days per ton, to at least 62, where we are using it now, with the provision that it would be nice to make some improvements in checks and things with data that are being generated in Japan and France.

And so we have agreements in place for 1 these data, and we have a Reg Guide that may be 2 3 revised to take all of this into account, but it is at 4 this point that the schedule kind of breaks down 5 because I really don't know exactly how we are going to wrap this up. 6 But I think in essence it is kind of 7 wrapped up and maybe Dana has another perspective. 8 Well, I like what you 9 CHAIRMAN POWERS: have said here better than I did on your first slide 10 on the nine topics, because you said it was resolved 11 12 there. MR. MEYER: Oh. 13 CHAIRMAN POWERS: And the experts got in 14 and looked at the thin little data that we have, and 15 said, gee, the biggest changes in 1465 actually come 16 17 because since 1465 was written, we have understanding of fission product behaviors, and I 18 would characterize that most of their changes is being 19 changes to the fission product phenomenology, rather 20 21 than the high burn-up effects if characterize them. 22 The database says, gosh, volatile fission 23 products look like they are released a little faster 24

from high burn-up fuel than from medium burn-up fuel,

but since they are high percent released, and 1465 is 1 an integral release, maybe it changes the timing and 2 there are some relatively minimal changes in timing 3 4 were made. And the more significant observation has 5 come out of the PHEBUS program where we are seeing a 6 7 lot more molybdenum release than we had in previous tests at any burn-up, and that seems to only get worse 8 as you go up in burn-up. 9 And those are the big changes that I 10 recall on this sort of thing. The database is thin, 11 and there are lots of things that we don't understand. 12 The VERCORS data and the PHEBUS data don't really 13 agree entirely on some fission products, notably 14 15 barium. There are lots of things, but to your 16 1465 isn't completely conclusion that 17 orthogonal to high burn up fuel is probably a pretty 18 fair assessment of the situation. 19 20 MR. MEYER: Okay. And the final issue that I want to mention is the one having to do with 21 fuel behavior during dry storage and transportation. 22 In the original plan, we said this is something for 23 the future that we don't have to worry about. 24 Well, the future arrived since then, and 25

there is now a need to renew some cask licenses and 1 license some new casks for burn-ups that are higher 2 than the 45 gigawatt day per ton that I believe was 3 the previous limit that had been approved. 4 We actually started working on dry storage 5 and transportation issues with some Surry fuel that 6 was medium burn-up fuel that had been in storage in a 7 demonstration out in Idaho for the last 15 years. 8 And as soon as that work came out of the 9 creep furnaces up at Argonne, in July of this year, we 10 inserted some creep specimens from the Robinson rods, 11 which are high burn-up PWR rods. 12 So at this time, we have high burn-up fuel 13 rods sitting in the creep furnaces, and these tests go 14 for something on the order of six months, nine months, 15 16 to a year. And during the next year we will also do 17 the isotopes measurements and other things that are 18 So this is a fairly short range effort that needed. 19 will give us a chunk of data that are needed for the 20 cask licensing and we will probably deliver that in a 21 research information letter to NMSS in 2004. 22 Now, there are a number of other factors 23 that affect their guidance, their review guidance 24 documents, and so it is not clear at this time whether 25

1	they are going to immediately make some revision, or
2	just hold that information until they are ready to
3	make other changes.
4	But I believe the research part of this
5	will be done in one year. So with that, I would like
6	to stop, and
7	DR, LEITCH: I just have one question back
8	on the reactivity-initiated accidents. Did I
9	understand you to say that you were making the
10	coolable geometry limit co-incident with the cladding
11	limits?
12	MR. MEYER: Yes, you did.
13	DR, LEITCH: I missed exactly what you
14	said in that.
15	MR. MEYER: That's exactly right.
16	MEMBER LEITCH: And what was the rationale
17	for that?
18	MR. MEYER: And the rationale for that was
19	that at low burn-up or zero burn-up, where the
20	original criteria were developed, and where we had two
21	different criteria, you did not have a mechanism for
22	dispersing or expelling fuels out of a cladding split
23	until you got up to a higher temperature, where you
24	started getting incipient melting in the fuel pellets.
25	And at that time, you now had a mechanism

for dispersing fuel, and breaking up the fuel rod 1 itself, and so we started out with this two-tiered 2 3 structure. 4 We end up with a situation where the high burn-up fuel 5 has a built-in mechanism for dispersing fuel, and as soon as you open up the cladding, it can cut fuel out 6 7 of the opening. MEMBER LEITCH: Okay. Yes, I understand. 8 MR. MEYER: Okay. So, Harold Scott is now 9 going to tell you about some of the work at Argonne, 10 and some of it is very recent, and I think you will 11 12 find it quite interesting. Let me just emphasize some 13 MR. SCOTT: topics that I am going to want to emphasize as we go 14 through in these experiments at Argonne in the hot 15 16 We did see a fuel loss, and I will show you some pictures and they will be in your handout of the 17 balloon fuel rods and the fuel actually -- and little 18 pieces of fuel coming out through that burst. 19 20 The other thing that we have observed in 21 two tests that were done with the radiated rods is that we get the same approximate shape and size of the 22 balloon and the burst. 23 At one point, we thought, well, this is 24 25 irradiated cladding, and it has got oxidation on it,

and it may give little tiny bursts, and maybe just little cracks. It will look a lot different. But so far we get sort of the same kind of balloons and cracks that we got previously. And there is some data on irradiated wouldn't be able to get down and make much of a balloon.

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

The Germans ran some tests 20 years ago and so we are not completely blind in terms of irradiated tubing. We also thought that with the high burn-up and the fact that the gap is closed that maybe the gas flow would be different, and therefore, maybe you wouldn't balloon, because the gas in the plenum

Or as soon as it ballooned a little bit, the pressure difference would go away, but it seems that the gas communication was at least good enough to give us balloons, and I will show you some of that information.

We also find that the rupture temperature was about what we expected, and so when we say, okay, let's put a certain amount of pressure on a rod, it balloons and bursts at about the temperature on the way up that we would have assumed from the information previously.

Okay. This is our largest program from a

215 1 financial standpoint, and so the boss, he spends most 2 of his money on my program here. Yuan Yan is the 3 principal scientist on this program, and Dr. Billone 4 is the principal investigator. 5 mentioned before in previous 6 meetings, EPRI was instrumental in getting these 7 limerick rods and the Robinson rods. These are Lee 8 test assembly rods, and the limerick rods are 9-by-9

The Robinson rods were made by Framatome and its predecessor, Siemens, and they are 15-by-15 rods. And as I mentioned before, they do have this typed fuel cladding bond and they might even be in some of the cuts.

It looks like the cladding and the fuel are sort of stuck together and they just don't fall out like they would as a bunch of pellets in a new cladding. So these are the kinds of effects that we would expect to be looking at.

So now I will talk about the kinds of effects that we are going to have. The main item in these oxidation kinetic studies was that the question had to do with whether the corrosion layer on the cladding to start with would make some difference in the following high temperature steam corrosion.

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

design.

1 And whether, of course, as the previous 2 slide showed, whether the fluence had some effect. So 3 this 1204C is the 2200 degrees F, and we are doing the same kind of experiments primarily that they did --4 that Cathcart and Pawel did, and other people have 5 6 done over the years a long time ago. 7 You oxidize them, and then you go in and measure these thicknesses. 8

Then we have the LOCA tests, the integral tests, which are sort of unique and new, and some of these tests have never been done before when we go through the whole sequence, because once again we have taken the rod and cut a piece out of it, and didn't really disturb the pellets in the middle of the section.

So we are looking again at this criteria. This equivalent cladding reacted is just a simple function of the weight gain, divided by the clad thickness. The trick of course is what is a clad As it gets thin, and as the rod balloons, thickness. you need to take that into account, and that is the way that this is defined on how to do that.

Ralph already showed you these little pictures of how we are going to do the bend and ring compression tests.

> DR. SIEBER: I have a question about the

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

1	details. Robinson fuels, 15-by-15?
2	MR. SCOTT: Yes.
3	DR. SIEBER: And it came from Framatome,
4	who evolved from Siemens, and Siemens evolved from
5	Exxon, right?
6	MR. SCOTT: Yes, and this was probably
7	made by Exxon in
8	DR. SIEBER: Now, Exxon autoclaved their
9	clad, which is a different process than General
10	Electric and Westinghouse, and a bunch of others, to
11	try and reduce the surface oxidation.
12	Did you find a benefit from that
13	autoclaving, and do you think that the fact that that
14	fuel clad was manufactured differently than other
15	brands that it had an impact on the data that you
16	have?
17	MR. SCOTT: Well, I think I show what
18	was the oxide thickness? It was almost a hundred.
19	Didn't we have that?
20	DR. SIEBER: That's pretty bad.
21	MR. SCOTT: Yes, and so we have these
22	rods had like many cycles, and so they started out and
23	were in a couple of cycles, and they took them out,
24	and they reconstituted them, and put them back in. So
25	the fact that it may have been sort of protected

1	cladding for an extent, the fact that they were in
2	there a long time to get up to this burn-up, because
3	they weren't particularly high enrichments, they had
4	to reload them into the assembly with other driver
5	rods.
6	DR. SIEBER: Right.
7	MR. SCOTT: So that they could reach that
8	and once again the maximum amount is this hundred. So
9	, some of them may have 80 and some may have 90. I
10	don't really know whether anybody said that
11	autoclaving technique helped these or not.
12	DR. SIEBER: Well, could I conclude that
13	it makes no difference as far as the data that you are
14	
15	MR. SCOTT: You mean is this oxide a
16	little bit different maybe than some other oxide?
17	DR. SIEBER: Yes. Could I draw that
18	conclusion and then we could just move on, or is there
19	a difference and did you look at it?
20	MR. SCOTT: I don't think we looked at
21	that.
22	DR. SIEBER: Okay. So let's just move on
23	then.
24	MR. SCOTT: Okay. I'm up to some of the
25	results here on

1	MEMBER FORD: And so the same point if you
2	are going to oxidize the inside. I am assuming that
3	Limerick rods are not barrier fuel?
4	MR. SCOTT: They are barrier fuel, yes.
5	MEMBER FORD: So the inside is severely
6	oxidized aren't they? When you blow steam through the
7	- <del></del>
8	MR. SCOTT: Well, no, for these oxidation
9	kinetics tests, they are OD oxidation only.
10	MEMBER FORD: Oh, okay.
11	MR. SCOTT: So we put them in the furnace,
12	and we plug up the ends so that nothing really and
13	we removed the fuel, and so we are just taking mineral
14	specimens that don't have any and we are only doing
15	the oxidation.
16	So there was no difference in the weight
17	gain between the unrated and rated. I will show you
18	a graph in a moment that shows this comparison for the
19	unrated. The Russians in our joint program with them
20	are working on this alloy, and it has niobium in it,
21	as do these two here.
22	And it turns out that Cathcart-Pawel does
23	pretty well on all of these alloys.
24	DR. SIEBER: Now, the Robinson fuel is
25	Zirc-4?
	, 4

wasn't really low-tin, but it was not the regular 1.6
old. It was maybe 1.5 or some other number. It was
a better version, or maybe selected from an ingot that
was a little bit because at that point they
realized that the more they got the tin down, the
better off they were.
DR. SIEBER: Yes, it had fewer car bumpers
in it.
MR. MEYER: Now let me show up
CHAIRMAN POWERS: What you are saying is
that the low level alloy agents don't make very much
difference in these oxidation kinetics, right?
MR. SCOTT: That's what we seem to find.
CHAIRMAN POWERS: That's terrific.
CHAIRMAN POWERS: That's terrific.  MR. SCOTT: Okay. Here is an
MR. SCOTT: Okay. Here is an
MR. SCOTT: Okay. Here is an unirradiated, and this is what we have dated all over
MR. SCOTT: Okay. Here is an unirradiated, and this is what we have dated all over the place with this kind of stuff, but not too much
MR. SCOTT: Okay. Here is an unirradiated, and this is what we have dated all over the place with this kind of stuff, but not too much data now for the radiated.
MR. SCOTT: Okay. Here is an unirradiated, and this is what we have dated all over the place with this kind of stuff, but not too much data now for the radiated.  But this is about and here is a scale
MR. SCOTT: Okay. Here is an unirradiated, and this is what we have dated all over the place with this kind of stuff, but not too much data now for the radiated.  But this is about and here is a scale under this 250, and so this is about 120, and this is
MR. SCOTT: Okay. Here is an unirradiated, and this is what we have dated all over the place with this kind of stuff, but not too much data now for the radiated.  But this is about and here is a scale under this 250, and so this is about 120, and this is it is hard to see the edge here, but this outer

1 Limerick fuel. 2 Okay. Thank you. DR. SIEBER: 3 MR. SCOTT: So I don't think you can see 4 it in here, but the barrier would be on this side, and it had maybe 10 or 20 microns of original corrosion on 5 the outside, and so in this case here for the 6 7 firradiated, you can't see it here, and I don't think 8 you can even distinguish it metallographically. 9 But this outside corrosion layer would be 10 here, and then the steam oxidation would have kept 11 eating away the zircaloy and forming zirc oxide at 12 this point. 13 And you have a nice boundary here for 14 these unirradiated, but in the irradiated, it is going 15 to be sort of tough to measure this thickness. 16 DR. SIEBER: That's your fault. 17 MR. SCOTT: That's how you get this. 18 call this prior beta because the temperature here was up to 2200 F. a long time, and this material changed 19 20 phase, and then as it cools back down, it comes back 21 maybe slightly different HCP than it was originally. 22 And there is some oxygen in here -- and I 23 will talk a little bit more about the movement of 24 that. 25 I think we ought to emphasize MR. MEYER:

1	that we have not done the oxidation measurements on
2	the Robinson fuel yet. So in everything that Harold
3	is going to talk about, we don't yet have a high burn-
4	up fuel that was heavily corroded in this database.
5	That will be started this fall, I believe.
6	DR. SIEBER: All right.
7	MR. SCOTT: That is a good point. This
8	data point would be, say, five minutes at the and
9	here is one at maybe about 10 minutes, and here is 20
10	minutes. So it seems to fit.
11	They took the old data out of the
12	Cathcart-Pawel report, and put it here, and then this
13	is like in cell, and so this is irradiated, and these
14	are unirradiated archive specimens from the GE
15	cladding.
16	This is some other material that we had in
17	hand in Zirc-4, hut it is unirradiated. So it is just
18	these ones that have an i (phonetic) that are the
19	irradiated.
20	CHAIRMAN POWERS: I am fascinated by your
21	vertical access. It says measured weight gain from
22	metallography.
23	MR. SCOTT: How do I get that? Okay.
24	Well, I can measure back on that graph that I had

before, and I go in here and I measure this thickness,

1	and I compute the weight gain from that thickness.
2	CHAIRMAN POWERS: When I look at that
3	layer and cross-section, it looks like the it is
4	topographic in its nature. And so it kind of takes a
5	measurement of the thickness. I mean, do you measure
6	it 50 times and take an average?
7	MR. SCOTT: Yes.
8	CHAIRMAN POWERS: Okay.
9	MR. SCOTT: And there is some and this
10	is Zr-02, and there is weight gain from this layer,
11	and this layer is maybe Zr-0.1, or .15 or something
12	like that. And there is a concentration grading
13	across it a little bit.
14	MR. MEYER: Tell them of the several
15	methods that are used to get the weight gain, because
16	that is not the only method that the lab uses.
17	MR. SCOTT: , Besides just weighing it at
18	some point.
19	MR. MEYER: They weigh it.
20	MR. SCOTT: Okay. Remind me a little bit.
21	MR. MEYER: Well, they have like three
22	different methods. The third one is escaping me at
23	the moment, but they do weight measurements, but this
24	metallographic technique turns out to be the most
25	accurate, and here they have done repeated checks to

1 see what their error has been, and this technique 2 seems to give them the best accuracy of all, and so 3 that is the one that we chose for this plot. 4 MR. SCOTT: And we get rather uniform --5 some of the tests that we did early on, this was a 6 variable, and we decided that there was something 7 wrong with the test, because most people who have done these kind of tests all the way around, you should get 8 uniform thickness. 9 The temperature in the furnace, we know it 10 11 is uniform all the way around. Here is a plot now that compares with the Cathcart-Pawel correlation, and 12 13 these different alloys. So what we did is we said 14 okay, compute at this temperature with the Cathcart-15 Pawel model, and it would then give a number right 16 along this line here. 17 So here is Baker-Just, and as we know it 18 gives more oxidation. The Leistikow is this one, and 19 this is the other one that probably has world 20 acceptance, in addition to Cathcart-Pawel. 21 The Urganic measurements were primarily 22 high temperature measurements, but they did enough 23 measurements at lower temperatures that they have a correlation. 24 This is the GE cladding. 25 These are

unirradiated, of course. We did some at Argonne, and 1 then we gave some of the same specimens to France and 2 they did them. So here is their answer here, and here 3 is our answer here, and so the point about this is 4 that it is sort of the same specimen, but done 5 slightly different. 6 They did double-sided oxidation, and they 7 have a different technique maybe of getting the answer 8 than we do, and so you can see the variability, 9 talking about some variability for the same duplicate 10 specimens. 11 So the fact that these lines and these 12 points don't all come down there, you are not going to 13 expect that because here is two materials that are 14 exactly the same. 15 This is M-5 data from the literature, and 16 this is not anything that we have measured yet, but we 17 expect to, and once again then this is the Russian 18 data that our colleagues in Moscow have -- well, here 19 is one here, plus one over there. 20 So it looked to me from this type of 21 figure that all these alloys -- we don't have any ZIRO 22 on here, but they are going to give -- you know, 23 Cathcart-Pawel can be used for all of those. 24 This is now my last slide on the 25

kinetics, and I wanted to go back I think to my figure, but now I don't remember why. Oh, okay. This item here about why does there seem to be some difference here on the cooling, or on the heating and cooling rates.

If the old data were taken with fast heatup, fast quench, but in these experiments, since we cool them down slowly, this layer, its thickness can change depending upon how fast or how your cooling down this material, and we cool that rather slowly.

Which is the case in the LOCA. I mean, it goes up to its peak, and it may not get to 2200 F., but it takes several hundred seconds sometimes to get back down to 800 Centigrade. Now we will go on to the integral tests, because these are the ones that we have had the most interest in.

The main idea of doing the oxidation test was to make sure that if it turned out that for irradiated material that Cathcart-Pawel was not the right number, or was not the right correlation then we would have to have a unique correlation to do these, because the idea is to oxide these specimens in the LOCA such that we get close to this 17 percent criteria or something like that.

And we have to include the prior corrosion

thickness. So these specimens are about a foot long, 1 and once again this is the 2200 F. number that we are 2 3 going to shoot for. This takes about 3 minutes to go 4 from --5 CHAIRMAN POWERS: Your next line, temperature ramps relevant to small-break and large-6 7 break LOCAs, is one that I am interested in. When you 8 say that those ramps are relevant to those particular accidents, presumably for which you calculate for 9 10 those rather, is it the -- what usually gets shown in connection with a small-break or a large-break LOCA is 11 the maximum temperature at any point in the core as a 12 function of time. 13 Did you use a ramp appropriate for a 14 15 particular rod in a core as a function of time? These kind of calculations MR. SCOTT: 16 would be done -- this is now during the -- not during 17 this initial CHF blowdown type thing, but later on, at 18 19 least for the large-break LOCA. 20 We looked at data and the calculations. So sort of like the hot-rod or --21 22 CHAIRMAN POWERS: Ιf you look particular rod, or did you look at the temperature, 23 24 which is the hottest location in the core, which 25 changes as the transient goes on.

1 MR. SCOTT: I understand that, and I am 2 saying that we have used a variety so that not 3 everything that we l'ooked at was the maximum power rod 4 that was running up the fastest rate. We also looked at some rods that were 5 running up at a slower rate. 6 Now, it may turn out 7 that some rods might go slower than this, and if they don't go very high, and if they only go from 400 C. to 8 9 800 C., I don't really care what the rate was for 10 that. 11 CHAIRMAN POWERS: I am more interested in 12 the rod that goes up and kisses 2200 degrees F., and 13 drops back down, and then comes back up again. MR. SCOTT: 'I guess I haven't seen any 14 15 calculations that do that. 16 CHAIRMAN POWERS: There were some that 17 were presented to us some years ago showing exactly 18 that kind of behavior. I don't know how general that 19 is. 20 MR. SCOTT: So these are the main -- this 21 one here is -- let me just say that this is assuming 22 that some rod in the core that had its maximum gas 23 release, and was driven hard, could have a pressure of, say, 2900 psig in it, and so under LOCA conditions 24 of atmospheric pressure there might be a delta p that 25

high across that clad.

But for the GE BWRs, this number is maybe 1250 psi. This is now a picture of the test train and that we have one of these thermal couplers on here will be the one that is controlling the furnace lamps from.

There is a pressure transducer at the bottom, and one at the top. This distance between these spacers was like 18 inches, and I sort of want to point that out, because visually in your mind now, turn this up so that when or after it has gone through its temperature, its high temperature, I am going to put water in at the bottom here, and the water is going to sort of come up in here.

And at some point here, it will start to boil and bubble, and throw or drop this up, and then this part of the rod will quench. Now, these are the parameters that we were using for heating up and cooling down the rod.

And I thought I would for you thermal-hydraulic guys, don't try to take this number and think about the FLECHT correlation or something in inches per second, because we have a different flow area, and our idea was to just get some water in there and start to get it cooled down.

4, 41

And after it reaches this temperature, we 1 would push the button and the water comes in. 2 again, we are looking for this equivalent cladding 3 reacted, and it would make a difference of about --4 well, wall thinning as Ralph showed when you get steam 5 on the ID and hydrating, and we are going to see all 6 7 of those. So the first few tests that we are doing 8 go for five minutes, and if we go for more than five 9 minutes, we would get way above the 17 percent 10 oxidation criteria. 11 So we have three specimens, and so we are 12 doing in three different times, we are going to make 13 So we did this first these different experiments. 14 test in August. These are the in-cells, and so these 15 are our first in-cell BWR tests. 16 We did guite a few out-of-cell tests with 17 unirradiated material to check everything out, and we 18 can then do comparisons between exactly the same sort 19 20 of set-up. So this one was done without steam, the very first one. 21 The next one, we did this one then in 22 September, and in this one we have steam and we take 23 it up here, but then we don't put any water in. So it 24 Hust cools itself down. We turn off the furnaces at

1 | 800 C.

So the next test we will do, probably in November, will be the complete sequence. So when we do this test here, this will be the first one with fuel, irradiated fuel, and the fuel inside and undisturbed, and through the whole sequence of quench.

I am going to now go to some of the outof-cell tests, and then I will come back and talk some
more about these in-cell tests. So here is this test
sequence.

At room temperature, we do some permeability tests and I will show you some graphs for that. Then we raise it up to 300 C. and do some more. The steam comes on, and off we go for this 3 minute ramp up to the temperature, and we are holding up here for 5 minutes.

But later on, we will do some longer tests, and it then cools down and we control the furnace to let it cool down at this rate. At this point then, the water in this next test that we haven't done yet is the one.

And it turns out from the out-of-cell tests that we have done that this sort of continues to come on down here. And even though we are adding water, and then all of a sudden it will drop down

1	pretty much like this. It really quenches quickly
2	once it gets to the point where there is enough water.
3	CHAIRMAN POWERS: Did I understand you
4	correctly that you have not done any quench up until
5	now?
6	MR. SCOTT: That's right. Well, out of
7	pile we have. We have sort of two set-ups. One set-
8	up is right outside the hot cell that uses
9	unirradiated tubing.
10	We have done all these sequences with
11	quenching with water, but not in-cell yet. So we have
12	done (a) and (b), and we are going to do (c). So here
13	again is the these 9-by-9s as I said are the GE-11
14	design with with the liner.
15	This is once again well, we chose this
16	1250 psig to try to give us a ballooning at about a
17	temperature that would give us a large balloon if we
18	could.
19	And if you go back to the old Oak Ridge
20	burst curve, they had a temperature and engineering
21	hoop stress, and there was maybe a high ramp rate
22	curve on there.
23	So this number turns out to be like 9.5
24	ksi. So you could look up on and that is what I
25	mean before about if I go back to that old Oak Ridge

unirradiated curve, and look at the KSI, and come over 1 2 to the burst temperature, and I see it would maybe be 3 750, lo and behold this test gave me about the same 4 number. 5 And once again we are getting 50 percent strain for this one, and this one is maybe a five 6 7 Now, this doesn't mean that the balloon was inch. this big for five inches. 8 9 It just means that if I look along the 10 profile, you begin to see a diameter increase over 11 this distance. So I will come to that picture in a 12 moment. But these are two more of the out-of-cell tests. 13 14 For instance, this one here would sort of 15 be the equivalent to this Phase A test that we did in-16 cells, since there was no steam involved with that 17 one. And this is part of our idea for deciding how to 18 get started with some of the experiments. 19 So here is a picture of this number three, 20 and I think in one of these viewgraphs before it 21 talked about a dog bone shaped burst. So that is what 22 they meant, the fact that it looked sort of a little bit like dumbbells at the end. 23 And this seems to be -- this may have sort 24

of collapsed back down in, and I can't tell from the

1	way that looks. And if you look and see, it was for
2	10 minutes. So this one has really high equipment
3	clad reacted.
4	You would almost have expected this one to
5	well, if we did a ring test on it, it ought to just
6	crack in little pieces. But it did survive cooling
7	down and handling.
8	Now, the next one did not happen and it
9	broke. Do you want a colored picture, Med, of the one
10	that I just showed you?
11	CHAIRMAN POWERS: The zirconium oxide
12	material is white and the stoichiometric material is
13	black, and color doesn't help very much.
14	MR. SCOTT: Okay. This is one that
15	survived the quench, but later then broke. The story
16	is that the guys were all done and they went off to
17	lunch, and when they came back, it had cracked apart.
18	CHAIRMAN POWERS: Well, they should have
19	taken it to lunch was the problem.
20	MR. SCOTT: It maybe better on your
21	handouts, but this is a shadow down here. They have
22	got a light up here, and so this is a shadow that you
23	are looking at here.
24	These thermal-couples are a little you
25	can't see it on here, but there is a little spot here.

1 They stayed on during the experiment and during the cool down and everything, and then just came off when 2 3 they handled it, the same way this one over here came 4 off. 5 And this one tested -- and this one again 6 had a high --7 MEMBER ROSEN: This is the one that I want to see the color on. 8 9 MR. SCOTT: I think I have that one, but 10 I don't know what I did with it. 11 MEMBER ROSEN: There is color on the 12 screen. I didn't bring one of those 13 MR. SCOTT: 14 with me. Now let me go off to these burn-up ones. 15 This is a fuel mid-plane, and so I am thinking high up 16 in the rod here, and then another one. This one is 17 going to be maybe between .8 and 1.2, and 1.1 space. 18 And we have seven of these rods, seven 12-19 What they did was that they shipped the rods 20 from the Limerick reactor out to Valecido, and they 21 come them into little pieces for us, and then we got 22 back all those segments. 23 And so we have a number of segments to look at, and part of the idea is -- and maybe it is 24 25 not so much in Limerick, but when we get around to the

Robinson rods, the higher up you are on the rod, the 1 more oxidation you have. So that may be a factor with 2 3 Robinson as to what grid span the sample came from. CHAIRMAN POWERS: Just a question on 4 5 nomenclature here. This Phase A, B, and C that you have under Limerick has nothing to do with the Region 6 A, B, and C, and your heating cycles? You previously 7 showed us a chart of your LOCA integral test sequence, 8 9 and you have an A sequence, and a B sequence, and a C 10 sequence. This one? 11 MR. SCOTT: CHAIRMAN POWERS: Yes. Those A, B, and 12 Cs, don't have anything to do with the A, B, and C 13 under Limerick? 14 MR. SCOTT: Yes. What I am saying is the 15 16 first Phase A test that we did, we went up to here, and stopped, and came back down in Argonne. The Phase 17 18 B test was one that went through this sequence, and this steam oxidized and came over here, but was not 19 20 quenched and just cooled down. 21 CHAIRMAN POWERS: Okay. The Phase C test that I am 22 MR. SCOTT: going to do in November is going to follow this path. 23 So A, B, and C all followed this part, but A stopped 24 25 here, and B stopped here, and C then follows the

whole.

CHAIRMAN POWERS: I understand now it does.

MR. SCOTT: Okay. So, yes. This is not October. This is the out-of-cell test. So here is the one where we were talking about it being a dogbone shape that we saw before.

I said before compare quite well with so far the two in-cell tests that we did. The shape looks the same, and the length looks the same, and the amount of strain is looking the same. So nothing seems to be out of the ordinary for these rods.

And we are hopeful that we will have to wait to do the Robinson rods. We won't be doing them until 2003. Now I am going to have some plots, and this first one and the next one are sort of at the beginning, and then later on just to show you what the pressure and the temperature do.

so we go up to -- this is the temperature up here, and so we are going to go up to 1200 C., and then come back down, and this would be this one here, right?

MR. MEYER: That is the temperature.

MR. SCOTT: Well, here is my ramp here at

1 the end here. Sorry. So, 300 C., and up to -- this 2 was this Phase A test that was in Argonne, and it stopped after it ballooned, and was turned off. 3 4 And this is a comparison between that in-5 cell test and an out-of-cell test. What we wanted to notice here is this in-cell test at the tail end, and 6 7 hotice how the pressure takes a while to fall down. 8 So what it means is that if I have large 9 delta p's, the gas can flow pretty well up and down in the rod. But if I get to a place where I don't have 10 much delta p, then the gas doesn't flow very well. 11 12 And we sort of expected that from the fact 13 that these are high burn-up rods. 14 MR. MEYER: May I say this differently, 15 because here is a case of looking at a glass that is half-full or half-empty. 16 I think the main thing to get from this slide is that the pressure took a nose-17 18 dive immediately in the in-cell test, just as it did 19 in the out-of-cell test. 20 Had there been a lot of axial 21 resistance, the pressure in the plenum would have fell off slowly, but it didn't, and so the gas is obviously 22 23 flowing easily from the plenum into the balloon area, 24 and depressurizing the whole rod quickly until you get 25 down to very low pressures, and then you begin to

1	notice this difference, because there is some flow
2	resistance, but it is not effective at the high
3	pressure differentials in really slowing down the
4	movement of gas from the plenum to the balloon.
5	DR. SIEBER: Now, the pressures that are
6	in there in a real situation would be the differential
7	pressure between internal rod pressure and the reactor
8	coolant system, correct?
9	MR. SCOTT: Yes.
10	MR. MEYER: That's
11	DR. SIEBER: And the pressure is
12	determined by the heating that is going on inside the
13	fuel element, and the cooling is taking place due to
14	ECCS or whatever else is taking the heat away. So you
15	aren't going to follow these curves. That the
16	phenomenon would occur as it is shown here in the
17	text; is that correct?
18	MR. MEYER: This is internal rod
19	pressure,a nd it would be like this is a real
20	situation, because you do have a plenum.
21	MR. SCOTT: At the top of the bundle
22	DR. SIEBER: Whether it bursts or not has
23	to do with the differential pressure across the board.
24	MR. MEYER: Correct, and we have that set
25	up about right, and this is showing that when it does

burst that the pressure in the plenum falls off very 1 quickly, which means that the gas is getting out from 2 3 the plenum and going through the fuel, and out the 4 opening. MR. SCOTT: Remember that in our case that 5 6 we are not too far away. The plenum is only a few 7 inches away from the burst, because we had data from Haldan for high burn-up rods when they changed gases, 8 and they do these kinds of experiments that show the 9 permeatability is rather low, and they try to measure 10 11 the hydraulic diameter of the gas, and it is almost 12 zero. And so we sort of thought maybe that the 13 gas can't flow very well. But as Ralph says, this 14 quickly depressurizes until -- and we don't think that 15 16 the ballooning is affected by gas flow, because we get this pressure change normally. 17 DR. SIEBER: Well, that is consistent with 18 your statement that the gaseous fission products are 19 20 distributed throughout the wall or the rod, and held 21 in the matrix end grain boundaries. But it does not require flow from the 22 plenum down to the point of ballooning the rupture in 23 24 order to get a rupture.

MR. SCOTT: Well, I don't know that the --

well, it is true that the -- well, okay. 1 2 DR. SIEBER: It doesn't require much gas. 3 But these in-cell tests, and MR. SCOTT: the boundary, and the outer rim of the fuel, it is 4 5 true that they have not been at this 2200 degrees F. I mean, the pellet rim region would be at 300 C., or 6 7 400 C., and not way up at 1200 C. DR. SIEBER: Right. 8 9 MR. SCOTT: So there is some -- I have put 10 a temperature transit on part of the fuel pellets, and 11 so there might be some gas release, but we didn't see 12 a big bump in the pressure. This is because the top 13 of the rod is heating up, and therefore the plenum is . 14 heating up. It is not from some gas release coming out 15 16 of the pellets. Now let me show you some of the 17 strains of these, and comparing the in-cell and the 18 out-of-cell. I may have marked on your handouts since 19 they are not in color, but this is the out-of-cell. 20 The zero degrees is where the balloon, and 21 so they turn the rod, let's say, with the balloon up, and then they measure how tall it is. Then for the 90 22 degrees, they turn it over 90 degrees and measure the 23 24 height once again at a difference.

So you can see that it has swollen some,

and all around the rod, and then these are the in-cell 1 ones and they once again come up to about the same 2 3 amount, and here is this nine degree one. So this one and this one sort of compare 4 and these two compare. And if this is 44 original, 5 half of that, 22, and 66, and so this is about a 50 6 And here is a good picture of how 7 percent swelling. it is now. 8 this 9 So at the bottom here, an unirradiated one, and it had like zirconium pellets or 10 something inside it just so we didn't have an empty 11 But you can see these little fuel particles in 12 there, and some of them fall out. 13 And once again these, because they were in 14 15 the reactor, I think this sort of reddish color is the color that these rods have because they have that 16 oxide layer, corrosion layer, on the outside. Here is 17 now a picture, and we can see the balloon itself up 18 19 close. This is the one that now went up burst and 20 no steam, and then was cooled back down. So that is 21 the kind of balloon and burst that we would expect for 22 a rod, whether it is radiated or unirradiated. 23 Now the thing that is also sort of new 24 that we didn't expect was this deposit. I don't have 25

1	the maybe I can go to the next one and then come
2	back to this one.
3	Before you saw that the fuel train was
4	inside of that quartz tube, and so here is the tube
5	again, and this is like a rag or something in the
6	background, and so forget that.
7	But here are these little fuel pellet
8	particles that have come out, and here is a black
9	deposit on the inside of this tube, and it turns out
10	that this is that inside the tube, this is where
11	the burst was.
12	So something came out of that burst and
13	pasted itself on the inside of that tube. We are
14	going to take that to a gamma scanning device, and see
15	if we can't see what it is.
16	You say a lot of moly comes out of these
17	fuel rods?
18	CHAIRMAN POWERS: That's at much higher
19	temperatures than what you have. You have not even
20	gotten close yet.
21	MR. SCOTT: Is cesium the only one that
22	would be sort of volatile at 2200 F.?
23	CHAIRMAN POWERS: Yes. These particles
24	may or may not be cesium.
25	MR. SCOTT: Well, okay. We will see what

1	they are.
2	CHAIRMAN POWERS: I remember that Dick
3	Laurentz, in his tests, reported in the burst test a
4	release of particulate and vapor cesium.
5	MR. SCOTT: These are the VI tests.
6	CHAIRMAN POWERS: No, these were tests
7	that he did on bursting rods many years ago.
8	MR. SCOTT: Oh, before that.
9	CHAIRMAN POWERS: But it turns out that
10	those things are extraordinarily important to the
11	transportation folks, because that's is their to
12	them that is the source term.
13	MR. SCOTT: So like I said, this is the
14	one that this is not a lot, less than a pellet's
15	worth of pieces. And now I am going to come back to
16	the burn-up case again here.
17	CHAIRMAN POWERS: I am going to face a
18	rebellion from my committee. I promised to allow them
19	to take a break and get some coffee before they close
20	downstairs. So could we take a 15 minute break here
21	and come back to the second test after that break.
22	MR. SCOTT: Okay. Thank you.
23	CHAIRMAN POWERS: It is a little bit of a
24	disruption to your presentation, but I think everybody

is following what you are doing pretty closely here.

MR. SCOTT: All right. 1 CHAIRMAN POWERS: So I will resume at a 2 3 guarter-of. (Whereupon, at 3:29 p.m., the meeting was 4 recessed, and resumed at 3:48 p.m.) 5 Let's come back into CHAIRMAN POWERS: 6 I remind everybody that Harold Scott is session. 7 discussing the LOCA tests, the first we've seen of 8 We've seen lots of plans, but not actual tests. 9 results. And Harold, I have to say that up to this 10 sense that overwhelming 11 point I've got the qualitatively not much has changed by going to the 12 higher radiation. 13 That seems to be from the MR. SCOTT: 14 information we've seen so far, but as we mentioned, 15 the Robinson rod was substantially thicker, corrosion 16 oxide may make a difference. We'll have to wait until 17 we get the first few tests from those. 18 CHAIRMAN POWERS: And I'll also say that 19 I remain concerned with exactly what you've got up 20 21 here nicely for me. You're a great straight man. With this heating schedule that you put up here 22 because my perception, rightly or wrongly, is that 23 this reflects the hot spot of the core kind of 24 analysis whereas especially for the Robinson test, I 25

think we're really interested in what an individually 1 rod perhaps at some point the hottest rod, but not 2 always the hottest rod, is actually experiencing, 3 which may not be monotonically heating up 4 monotonically cooling down, but rather going through 5 wild gyrations. 6 But remember that we don't 7 MR. SCOTT: really get any oxidation cooking until we get up into 8 9 here. CHAIRMAN POWERS: But remember, that 10 you've already got it oxidized. 11 MR. SCOTT: The outside is. 12 CHAIRMAN POWERS: The external oxide and 13 especially with Robinson fuel for your roughly 100 14 microns, it's getting very close to the point where 15 16 that oxide becomes very susceptible to thermal shock 17 induced spalling. MR. SCOTT: But also, the kind of data we 18 have from various ballooning and burst tests, this 19 isn't particularly too critical. This is 3 or 8. 20 21 It's going to come up here and I wouldn't think we'd get too much effect on -- what we will get some effect 22 on is the fact that this material is irradiated may 23 change this what we call alpha-beta transition 24 25 temperature which will depend upon -- Hee Chung says some of these bursts look like -- they're a little bit different than the ones he would have expected at that temperature because they've sort of crossed over into another crystallographic -- I put this up to remind you again because I'm going to show some vu-graphs. We're doing a permeability test, a gas flow test at -- down here and then at 300 C and this was the A one. Then the B one goes through the high temperature oxidation.

MR. MEYER: While you're changing slides, let me comment to Dana.

I think we will get the information that you're interested in from the integral tests where we will be looking at the oxidation level in the balloon region where it has deformed and broken up any heavy oxide that was in that region, so even though we don't jerk it up and down in temperature, there is going to be a big balloon deformation taking place that will mechanically shake up the oxide and where we will look in great detail.

MR. SCOTT: Yes, we haven't had a chance to do any metallography on these specimens yet. So this is the low temperature, this one and the next one showing you how well this upper pressure transducer and the lower pressure transducer track each other and

248 as we just mentioned before, Ralph reminded us that 1 this one tracks very well, right here at 2 So that's sort of what's critical, the 3 beginning. fact that it pales off here a bit here probably 4 doesn't have too much difference in the behavior. 5 So now we're going to go to Phase B which 6 steam when through the 5-minute oxidation. This was 7 the pattern here and then we ramped up to -- and I 8 think I've got some graphs here I may have shown 9

think I've got some graphs here I may have shown before. We got this little pressure peaking because

the plenum heats up again.

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

Here's the burst temperature. All of these, you'll notice, A, B and the out of pile, out of sale tests all had for this same gas pressure had about this 750 C. In fact, I think I noticed that before here. One of them -- they're the right order. If the pressure was a little bit higher, it -- the burst temperature was a little bit lower which is what you'd expect.

So how we're at the higher temperature permeability and it looks about the same. Once again, it's a little lag in this lower pressure transducer to see the pressurization, but then it quickly catches up. And I have the downlay side.

1323 RHODE ISLAND AVE., NW

WASHINGTON, D.C. 20005-3701

Now we're off to the test, the most recent

test we did in September. This is this little heat up 1 here where the pressure goes up. This is the bursting 2 3 here and then it goes on up and starts to oxidize. If you wonder why we don't MR. MEYER: 4 transducer at high 5 show the lower pressure temperatures, it always fails. We've got -- the steam 6 is doing it in and we have to do something about that. 7 In the first test, it didn't MR. SCOTT: 8 hardly work at all and the second test, it works, but 9 wasn't reliable. 10 Once again, here's this 50 percent strain. 11 gone back and subtracted out oxide haven't 12 thickness, so these numbers will in the report might 13 be slightly different and once again, the shape of the 14 burst opening was sort of like what we saw in the 15 underrated experiments. 16 Here's now a plot of those. I was just 17 talking during the break, Robbie Montgomery, his code 18 will calculate, he can see this shape here. My code 19 doesn't do that. I get this number and this number, 20 but I don't -- I'm not able to calculate the shape of 21 22 this. We don't know how you do it, 23 actually, we do. 24 Okay, now I'm up to a picture here of 25

1	these of the one I just showed you here, the
2	ballooning and you can see this is this zero
3	degree, if I measure from here up to here, that's the
4	so-called zero degree. If I roll it over, then I get
5	a 90-degree measurement here and this is when I said
6	before like this is the from maybe here to here is the
7	amount of ballooning that I get.
8	And part of this point is before when
9	Ralph showed you that schematic about how's he going
10	to cut a ring near the balloon, but it's still got
11	that high hydrant, so we think up in here, there's
12	going to be, you can get rings up here that we don't
13	think will have much hydrant because it's not much
14	weight for that extra steam and hydrogen to get all
15	the way up here. But in this part here, we can get
16	several of these rings out of here and then we can say
17	from here over to here and do this 4 point bin test.
18	CHAIRMAN POWERS: I take it you are
19	lobbying heavily to convince Argonne that they ought
20	to become metric?
21	MR. SCOTT: Weren't all these metric,
22	centimeters and C.?
23	MEMBER FORD: An inch?
24	(Laughter.)
25	CHAIRMAN POWERS: You can explain to him

these modern measurements. 1 MR. MEYER: The hot cell was built in the 2 This ruler has been in there every since. 3 (Laughter.) 4 It's probably a little over DR. SIEBER: 5 waste by now. 6 MR. SCOTT: Okay, I mentioned before this 7 dark deposit on the tube that occurred. The other 8 question we sort of had was does that deposit now 9 affect the temperature behind it since the lamp was 10 trying to send entry through and I was told that they 11 were going to check that out and do some tests with --12 put a device inside of that tube that has a deposit 13 and see if it can -- if that shadow actually makes any 14 difference. 15 But we were told it was rather thin, so I 16 think it's not going to make much difference. 17 again, we're talking about the amount of fuel that 18 came out. We put a little basket at the box so we can 19 catch the fuel if it falls out during the test and 20 then later if when they take this thing over to some 21 table, but they try to keep the brake up so nothing 22 else falls out. But a pellet is maybe 10 grams. 23 We're only getting maybe half. 24 really able -- I've not seen these and you can't see

it in one of these pictures here. I'm going to show a little bottle. Here's these pieces. They're small, but I don't know if they look like shards or if they -- the size of a bb size, but we'll characterize those.

The other thing that came up is in preparation of these specimens are we doing anything to the fuel rod and the pellets inside that maybe would make a difference in the answer. Is our experimental technique affecting our answer? So we have these specimens that we cut and we can look in there and say okay, if I drill out the top of the rod to put a little plenum in it and a cap on it, have I somehow vibrated and cracked the pellets six inches away, so we're going to do some -- and so far we don't see that. We can head it off and it looks just normal.

Once again, this dark deposit, to see if we can see what it is. We'll calculate more exactly the equivalent cladding reacted and we'll look at the -- we have a hydrogen determinator device that if you take a specimen, we'll be able to see what the PPM to hydrogen was at the various locations.

Then we have the -- as I said before, we've done these out of sale tests with the quench.

We'll take that quench system and put it in the cell 1 so in November we can do this phase C test that's the 2 full LOCA sequence. And that's the end of mine. 3 Are there any questions? 4 CHAIRMAN POWERS: Any other questions for 5 Harold? 6 I guess I just want to MEMBER LEITCH: 7 make sure I'm coming away with the right conclusion 8 I guess what I'm hearing as far as this high 9 burnup fuel is concerned from the work that you've 10 done so far, and I know you're anxious to see the 11 results of the Robinson test, but from the results of 12 high burnup fuel at Limerick, we don't see that much 13 unexpected or different than you would have expected. 14 Is that a correct assessment? 15 let me answer that and say MR. MEYER: 16 that's a correct interpretation of what we showed you, 17 but keep in mind that what we're looking at are 18 embrittlement criteria and evaluation models and we've 19 now looked at data relevant to three of the evaluation 20 kinetics, the rupture oxidation models, the 21 temperature pressure conditions and the ballooning 22 strain. 23 Those three at least on the low corroded 24 BWR fuel look unaffected, qualitatively unaffected by 25

1	burnup. We have not get looked at the embrittlement
2	which is the one that is most likely to be affected by
3	burnup because it should be directly affected by
4	hydrogen absorption and there's going to be more
5	hydrogen in the burnup specimens than in the fresh
6	fuel.
7	So we've looked at three important models,
8	but not at the criteria.
9	MEMBER LEITCH: Then, of course, the
10	Robinson work will be very interesting because they
11	have thicker oxidation.
12	MR. SCOTT: And it's hydrogen levels are
13	substantial. When you have 80, 90 microns of oxide,
14	you get substantial hydrogen.
15	MEMBER LEITCH: Yes, okay, thank you.
16	MR. SCOTT: We had a paper that Argonne
17	issued, about 10 pages, back in June. It's in ADAMS.
18	I gave Med a copy if anybody wants to get it. It sort
19	of shows some of the graphs I've showed and describes
20	more details of these ECR calculations and the fact
21	that we get sort of a similar oxidation for the
22	different alloys.
23	MR. MEYER: So now I'm going to come back
24	to the RIA and ATWS situations and just hopefully
25	demonstrate a little bit of progress in the last year,

but I don't think we'll reach too many conclusions 1 that you haven't heard before. 2 I'd like to summarize the pulse width 3 situation to show this Vitanza correlation and then to 4 describe briefly the method for making temperature 5 corrections. 6 This is just two typical cases that were 7 run with the PARCs 3D neutron kinetics code for rod 8 worths that are reasonable, about \$1.20. And in fact, 9 they produce relatively low energy pulses. This is a 10 plot of the power for these, a beginning of side and 11 an end of cycle. They're different. The calculation 12 takes the plutonium build up into account and other 13 14 things. And here is the enthalpy, the fuel pellet 15 enthalpy for those two cases and you can see, indeed, 16 that the enthalpy peaks rather slowly and it's a low 17 value on the order of 30 to 35 calories per gram, but 18 it started at 18 calories per gram, so the increase 19 was only 15 to 20 calories per gram. 20 Now based on a fairly large number of 21 cases and I think you've seen this slide before, 22 Brookhaven has used that code to look at pulse width 23 as a function of the change in fuel pellet enthalpy 24 they've done that for a lot of different 25 and

assumptions.

2

1

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

Not on this slide, but on a similar slide, results from other people's been placed have calculations, from some of the vendor calculations and except for quibbling a little bit about the exact value, there has really never been any serious criticism of this finding. It also checks well with the Norhung-Fuchs equation, so there's analytical basis that doesn't rely on big codes and then there's big codes and there's other people's big codes. the bottom line is that if you have low energy pulses that are broad, if you have high energy pulses that are narrow, and this morning EPRI, talking about pulses that have pulse widths no greater than about 30 milliseconds and if you, I'm sorry, no less than about 30 milliseconds and if you look on the chart you will see that those energies then are all less than 30 calories per gram.

Now let's see what I have next. If you are interested in running a test at a low energy that is comparable to what you would predict for a PWR in this accident, then you would want to run a test at maybe 30 calories per gram and 30 milliseconds pulse width. But if you're interested in exploring the energy range where the cladding is going to fail,

1	pressurized water reactors, but the pulse width for
2	the boron dilution pulses look right in line with the
3	rod ejection pulses.
4	I'm not sure if this conclusion about the
5	boiling water reactors is well examined or but it
6	kind of makes sense that there could be a difference
7	and it sort of the characteristic of a core and
8	CHAIRMAN POWERS: Still, I think the way
9	you went about a decision to drop the explicit
10	consideration of the rod drop accident of the BWRs was
11	appropriate use of risk-informed decisionmaking,
12	guiding your research program.
13	MR. MEYER: You can do that or if we solve
14	the problem for the PWRs, then the BWR analysis
15	CHAIRMAN POWERS: It might like falling
16	off a log, right.
17	MR. MEYER: Right, right.
18	MEMBER LEITCH: In the BWR, have you given
19	credit for the velocity limiter or is this just an
20	instantaneous
21	MR. MEYER: No, no, no. The velocity
22	limiter is taken into account.
23	MEMBER LEITCH: Thank you.
24	MR. MEYER: I'm going to come back to
25	that. I didn't put the slides in the order that I

wanted to have them in. And I'll just talk a minute about the temperature effect related to pulse width.

Well, pulse width, we imagine has several effects. One is through temperature and one is through dynamic fission gas expansion. We have not done any examination of the dynamic fission gas expansion hypothesis and I don't think EPRI has modeled that. It's a hypothesis that's out there and it might account for some of the scatter in the data, but certainly, we ought to be able to handle the temperature effects and I just want to make a few simple comments about it. We haven't done it yet, but we're beginning to work on it.

Here are three results from three calculations that are kind of illustrative. The three cases that we took resembled NSRR pulse, a PWR pulse and a CABRI pulse. All three of these pulses have a total fuel enthalpy of increase of about 100 calories per gram.

What we did was plotted cladding temperature as a function of fuel enthalpy, rather than temperature. And the picture to have in mind when thinking about this is we have a reactor that can give us 100 calories per gram fuel enthalpy increase and the cladding that's going to fail at 80 or 90

calories per gram. So we want to look in the range of 80 or 90 calories per gram. This is the time at which the cladding is going to let loose. It's going to fail and find out what the cladding temperature was at that time. And then try to relate that to some mechanical properties or something.

So here are the cladding temperatures. Now, the NSRR temperature is very low because it started low. It started at 25 degrees instead of 300 degrees Centigrade. Had it started at 300 Centigrade it would be very close to the 10 millisecond line.

And the 30 millisecond pulse at a given fuel enthalpy out in the range where you might expect failure is about 70 degrees too high.

Now one of the things that we think we notice from the data are that the total plastic strain in the case with the real broad pulse, the 30 millisecond pulse was a little less than the total plastic strain was in the 10 millisecond pulse. Now the fuel enthalpy was the same, so the pellet expansion should be the same and the difference that we think is there and I spoke to Rob about this earlier and we're not sure of it, but we'll look at it, is that the cladding is going to increase its diameter just from thermal expansion. And since it's

hotter, it's trying to run away from the pellet and it's able to run away a little bit better when it's hotter.

So I think there are really two effects to look at here. One is thermal expansion. The other is the mechanical properties, the temperature effect of the mechanical properties. And so here is a plot of a collection of data that we have that shows total elongation as a function of temperature almost in the right temperature range. It doesn't quite go high enough, but you see here exactly the same kind of spread that you saw in the CSED curve, because the CSED curve is a reflection of the total elongation measurements.

And so we will in trying to use this, we will experience exactly the same kind of difficulties that EPRI experiences with data like this, but you know, you can say the temperature effect is between zero and this and we can look at that parametrically.

From thermal expansion and from the tensile data, we can then get a strain increment that is related to the temperature difference. You could call it strain, I guess on the thermal expansion and then we can relate that to the enthalpy chain. So this is simply the -- for that 10 millisecond pulse,

the cladding strain is the function of enthalpy increase. And so we can convert the delta Ts to delta strains to delta Hs and then take them back to the paintbrush slide and move the data around and claim that we have made a correction for temperature, although we make no claim about any other effects like the dynamic fission gas or perhaps some pellet lock up or something like that.

So that's the method. Hopefully, you see a little progress from a year ago where we are. I'm going to try and go back now and pick up those other slides.

Okay, so I mentioned that we were also trying to use an empirical correlation and this is Vitanza's correlation and I only show this to indicate that the failure level in the correlation is dependent on a number of parameters, on the burnup, on the mechanical properties of the cladding, on the pulse width, on the oxide thickness and on the cladding wall thickness.

Vitanza compared his correlation with the failures in the CABRI data sets and there's one more point on this then. EPRI has showed this is the MOX data point, so he predicts the RepNa-8, RepNa-10 and I think it's the RepNa-7, the MOX failure quite well,

but like everybody else, can't predict RepNa-1. 1 2 And I agree with Rosa and EPRI that RepNa-3 1 is an outlier. I'm probably, less diplomatic and 4 more conclusive in my view because it's our contractor 5 who has said that the preconditioning temperature soak 6 has probably caused the embrittlement of this and has 7 written a number of detailed descriptions of his 8 observations of severed hydrides and all kinds of 9 things to support that position. 10 So I am inclined to believe Hee Chung from 11 Argonne National Laboratory that that is the main 12 reason that this test result is not reliable. The 13 other factors that Rosa mentioned are also legitimate 14 areas for looking into and I think this whole thing 15 will be wrapped up in another few months. Hopefully, 16 we'll get that behind us. 17 CHAIRMAN POWERS: I'm just not sure of the 18 Japanese data. 19 MR. MEYER: No, it doesn't. I don't 20 recall whether -- did he --21 MS. YANG: No, I think Carlo Vitanza only looked at high temperature data. He basically just 22 23 took the CABRI data and fed it into the equation that 24 you presented. 25 MR. MEYER: I don't think this correlation

is ready for service, but it is interesting. 1 2 moving in the right direction. I've spoken to Carlo 3 about it and he's not only willing, but eager to work 4 with us on this and I'm hoping that we can work with 5 him to develop this correlation a little more broadly. CHAIRMAN POWERS: Correlations 6 7 strictly empirical type like that suffer needlessly 8 when you try to extrapolate it and of course, you're 9 trapped in extrapolation here because you're doing 10 your tests in situations that people can find a litany 11 of fault where your data base is coming from. 12 A phenomenological understanding is always much better, but when I did experimental work I always 13 said well, let's get an empirical fit of the data 14 15 first and then we'll work on the phenomenological. Sometimes that didn't work out. So it may have some 16 17 virtue to it, a less desirable outcome than Hee Chung 18 talking about hot short metals and things like that. Well, as I mentioned this 19 MR. MEYER: 20 morning we really have a multiple approach to this, one of which is a code calculation which involves the 21 mechanical properties in a manner that's similar to 22 23 EPRI's. 24 I frankly think that in the end we don't

need exquisite accuracy on this thing because I think

we're going to have a margin of a factor of 2 on a failure limit that is clearly conservative. If you don't crack the cladding, you can't have bad things happen.

So if it works that way and we have some uncertainty in this correlation, in my opinion that would be tolerable.

Okay, on the BWR power oscillations, we talked to you a year ago about the implications that we drew from the PIRT elicitations and I have those on the next two slides and I don't intend to read through those. I just want to tell you about two new steps forward on this.

From the PIRT implications, there was a conclusion that the repeated power pulses would probably not cause PCMI failures and that in the end this would be a high temperature transient and that the temperature would be the damage mechanism. So what we have from Japan now are two tests in which they did repeated pulsing. And let me see if -- they used BWR rods, two of them, with modest burnup, so 25 gigawatt base to turn 56 and they found that the mechanical interaction didn't enhance, wasn't enhanced by cyclic loads which was one of the things we were worried about, sort of a ratchet effect.

268 This is a slide of their data which they 1 will be presenting at the Nuclear Safety Research 2 Conference in a couple of weeks and it shows the 3 cladding elongation which is pretty small, what am I 4 looking at here? The relative rod power is not on the 5 scale and then you have the temperature which is --6 ah, here is the cladding elongation and these are the 7 temperatures and this will be explained at the NSRC 8 conference in a couple of weeks. 9 The other thing that we have done is we've 10 agreement with STUK in Finland for signed an 11 They had a little cooperation in the analysis area. 12 thermal hydraulic code called GENFLO which they 13 coupled to FRAPTRAN and used that to try and analyze 14 the rather active feedback that goes on between the 15 hydraulic conditions and the fuel rod conditions in 16 this transient. 17 This code was actually installed out at 18 Battelle just almost a month ago now and we've run the 19

This code was actually installed out at Battelle just almost a month ago now and we've run the code on some sample cases and are going to plan our attack, our analytical attack in the next year.

So with that I think we're ready for Sud Basu who will talk about the fuel behavior under dry storage conditions.

CHAIRMAN POWERS: Thank you. Are there

20

21

22

23

24

1 any additional questions for Ralph on his final 2 presentation or anything that's gone 3 suppose? 4 Seeing none, we'll proceed. 5 MR. BASU: So at the end of the day I will 6 talk about some old stuff and I mean literally old 7 stuff. We'll talk about spent fuel rods which were in 8 the reactor for about three years. Then they had a residence time in a wet pool for another five years. 9 They were taken out. Went through vacuum drying and 10 11 they were stored in dry casks for about 15 years. 12 Back in 1999, they took some assemblies out, did some 13 observation on their behavior and that's what I'm 14 going to talk about. 15 The scope of the program is looking at the 16 post-storage and by that I mean 15 years of storage in 17 dry cask. When we took them out, post-storage 18 characterization of these spent fuel rods. I'm going 19 to actually focus more on the creep testing of fuel 20 rods and I'll touch upon --21 (Pause.) 22 I'm going to, as I said, emphasize the 23 creep testing of fuel rods and I'll make some comments 24 on the post-creep mechanical properties. 25

We are looking at and we have actually

looked at Surry fuel rods with a medium burnup, less 1 than 45 gigawatt day per metric ton. We are currently 2 looking at, we have started the campaign on high 3 4 burnup cladding. Ralph alluded to that. 5 The focus of this presentation is on Surry rods because we had results to share with you. 6 7 rods which we have actually sampled from the dry casks 8 have an actual burnup of 36 gigawatt day per metric 9 ton. As I said, they spent in wet pool for about five years and in dry storage since 1985. 10 11 Now why are we interested in this stuff, 12 this old stuff? The rods that are stored in dry casks 13 are the dry casks, actually the dry casks are coming up for license renewal as early as 2004, not all of 14 15 them, obviously, but the population of dry cask will 16 be coming up for license renewal. 17 MEMBER ROSEN: How long were they licensed for originally? 18 19 MR. BASU: Twenty years, original license 20 period is 20 years. They're coming up for license 21 They'll be submitting license renewal renewal. 22 application. About this time, they'll probably submit a couple of them and they'll be submitting more and 23 24 more and there's a two year period between the

application and the ramping of up renewed licenses or

what we call certificate of compliance.

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

In order to issue that certificate of compliance, we have to assure that these casks can go up to another 20 to say 100 years. That's the license And of course, in order to assure renewal period. that, we need to assure ourselves that the fuel rods which are stored in dry casks are in good condition to be restored. So that's the incentive that is driving the medium burnup work.

And there is the incentive for the high burnup work that's the new licensing. We want to be able to verify the validity of the efficacy of Part 72 rule and how that transfers to tech specs in the Spent So that's the incentive for Fuel Project Office. doing the high burnup creep studies that Ralph alluded to and I'll touch upon that as time permits.

Part 72 says that the spent fuel in dry casks must be protected from degradation that leads to gross ruptures. That's a very broad definition. That definition has been translated in the technical specification, if you will, of the staff guidance work as cladding that should not have or must not have more than 1 percent creep strain over the period of the life in dry cask.

It certainly must not have crumbling or

you know total loss of geometry, so to say. And then of course for -- we need to look into mechanical properties of these rods so that during restorage or transportation that these rods do not lose their geometry or do not lose their strength, so to say.

So we need the creep and mechanical properties data and that's what the focus of this, the work that I'm going to present. I'm going to go very quickly through the post-storage characterization part because that's kind of an uninteresting part in terms of observations that we made.

What we did was we took 12 rods from an assembly that we recovered from an open cask and we did the peripherometry of these 12 rods to see the diameter changes and what we found is that the diameter changes first of all are pretty uniform and Very little variation they're about .6 percent. azimuthally or axially and what that transfers to is a thermal creep during that 15 year of storage life to less than .1 percent. Very little. Very little.

Then what we did is we took 4 of these 12 rods and we did -- we punctured holes and we did some gas analysis, fission gas analysis using fission gas analysis -- well, laser puncture technique so that we can do fission gas analysis. What we found is that

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

fission gas release is about 1 to 41 percent which is well within the range that you would expect from these rods stored in dry casks for about 15 years. And of course that again translates to some internal gas pressure of 3.5, around that, which is then within the range, so that's why I said these are all uninteresting results and that's -- there's nothing exciting about what we found. It's all expected results. We did metallography. rods out of these four rods.

Not all four of these rods or not all four segments, but we chose two Again, these were so uniform in every respect that we didn't have any problem choosing any two rods from the inventory. chose two rods. We cut up segments and we did metallography and what we found is the rod thickness varies from 20 to 40 microns, about that.

The hydrogen content varies from 200 to 300 PPM. No hydride reorientation, not that we expected any hydride reorientation, but we wanted to be sure there's no hydride reorientation during the vacuum drying period and during the external storage period and that's what we found.

We did also some microhardness testing and what we found is the microhardness is about 240 DPH

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

1 which tells you that there is no annealing that took 2 place during the storage period. Again, nothing 3 unexpected. 4 MR. MEYER: What is DPH? 5 MR. BASU: DPH stands for Diamond 6 Perimeter Hardness. 7 MR. MEYER: Okay. 8 MR. BASU: That's a hardness testing 9 You use a diamond cone, diamond shape. 10 indent the surface and you see how much deformation of 11 the surface. That's what it is. 12 MR. MEYER: Thanks. 13 MR. BASU: As I said I'll be focusing more 14 on the creep test because that's what we really want 15 to know how much creep these rods have gone to in 15 , 16 years of dry storage life and how much residual creep 17 lies ahead. So we came up with the metrics and these 18 are seven tests. Five of these tests have already 19 been conducted. The two that you see at the bottom, 20 6 and 7, have not been done and I'll come to those in 21 a little while. 22 The conditions for the creep tests, the 23 conditions were selected to represent pretty much the 24 temperature that you would expect in the beginning of

life storage, dry storage of 360 to 400 degrees.

That's where these temperatures come from.

If you take that temperature and you try to run a creep test within a finite time frame, you're not going to see any creep whatsoever. So what we did, in order to do some creep studies, we had jacked up distress to about twice or a little more than what you would experience what these rods experience in the beginning of dry cask environment.

Again, the purpose of these creep tests are multifold. We want to, of course, know what is the residual creep life in these rods. Do they have 10 years left, 50 years left so that that will give us an idea of whether we can really renew the cask license, but of course, also to generate the primary and secondary creep data so that we can use the date to develop correlations or to verify correlations that are in the code and in the model.

I'm just going to go through very quickly because these are standard creep tests. There's not much to explain here. The 3-inch specimens, the cladding segments for the fuel and then refuel with zirc pellets and the specimens were pressurized with Argonne gas. The pressurization system has the capability to pressurize up to 6000 psi which translates to something on the order of 330 megaPascal

1	hoop stress. Okay?
2	Excuse me. My throat is drying up.
3	It's a fancy regulated system that Argonne
4	has which can regulate pressures up to Class 1 of 10
5	psi.
6	CHAIRMAN POWERS: I'm impressed.
7	MR. BASU: It is impressive.
8	Unfortunately, I don't have a picture to show you
9	here, but it looks fancy.
10	I have a picture to show for the specimens
11	loading in furnaces to do concurrent creep testing, so
12	we can do more than one at a time. By way of
13	measurements, we of course to the temperature, the
14	temperature and pressure measurements as the control
15	parameters and in terms of the measured parameters to
16	derive the strain and strain red. We did the diameter
17	measurements at multiple axial and azimuthal
18	locations. We also did length measurements to verify
19	whether or not there is anistropy in the creep
20	process.
21	Again, the dry data from the diameter
22	measurements of hoop strain and the strain rate and
23	strain time history
24	MEMBER FORD: Why did you put it in
25	zirconium pellets?

1 MR. BASU: Oh, this is to stimulate the pressure inside what is representative of what you 2 3 would expect if you were actually doing testing with 4 fuel inside. 5 Even though you filled it MEMBER FORD: 6 with argon -- pressurized it with argon? 7 MR. BASU: Yes, because some of the energy 8 will be absorbed in the pellet as opposed to putting all the energy to cladding. 9 That would not be 10 representative. 11 Okay, here is the 3-inch specimen that I 12 am talking about. This is a 3-inch specimen. It's 13 the end cap and this is the pressurization system, 14 welded to it. It's going to an argon chamber to 15 mitigate any contamination due to -- if this was an 16 air or open space there would be oxidation, perhaps, 17 so to mitigate that we have this chamber and this is 18 the creep system or the assembly of furnaces. 19 are smaller furnaces which can accommodate 1 sample 20 leech, and this is the largest furnace which can 21 accommodate 3 samples. You can pressurize these samples at different pressures, but of course, here, 22 23 the temperature for all three samples would be the 24 same.

the

photograph

here's

Okay,

25

the

of

diameter measurements using laser profilometry again. We have the spindle here. The sample is taken out directly from the furnace. Put in a spindle here and then you can actually move this axially in this direction to get axial measurements. You can also rotate this to get the azimuthal measurements at 20 degree increment. So you get, what is it, 18 measurements for each axial location and you get much more axial measurements.

Okay, so what do we get from that layer of performing measurements are these diametral data and what these circles, the perfect circles, if you will, show, showing are the diametral marker, one for 8 inch. One for 9, one for 20, etcetera, and these are the before diamond if you will, constructed from 20 azimuthal measurements.

And as you can see that kind of progress is the creep time progresses.

These are, of course, the average over the length of the segment, length of the specimen, and I'm going to show you what the variation of the length of the specimen, it's really not much. So we have a 10, 2, 3, 5, 7, 9 data points around the axial direction and what we do is we average the five middle ones and just discard the 10, 4, 2 on each side. So that's how

you generate the data that you saw in the previous 1 plot. 2 3 Okay, so those are in terms of measurement and some kind of data reduction and then we come up 4 5 with results in terms of what we can relate to for 6 creep and that's the average strain. That's in 7 percent. And the strain rate that we can construct 8 from average strain measurements. 9 There was no failure in all five tests 10 that we conducted. 11 CHAIRMAN POWERS: Yes, but did you take 12 No. 4 out quickly so it didn't? 13 MR. BASU: No. That's a good question. 14 What happened was we kind of tricked a little bit. Didn't mean to trick you guys. Within No. 3, that's 15 16 No. 3 at 400 degrees, 190 MPa for this length of time 17 and we saw an average strain of 1.03. Of course, no failure. And then we said really, that's a very small 18 strain and it's been a fairly long duration, so we 19 20 took that out, put it back. We jacked up the stress. 21 So what you see here for this duration, the additional 22 strain that you accumulated is this 5.83-1.03 or about 23 4.8 percent, by just jacking up that stress. If you want, it's 3A and 3B experiments. 24 25 So, all right?

1	MEMBER FORD: Those strain rates are the
2	average strain rates? Those strain rates are the
3	average?
4	MR. BASU: Yes, based on the average
5	strain.
6	MEMBER FORD: Because it's not a
7	logarithmic creep log, decreasing the time,
8	presumably.
9	MR. BASU: Yes.
10	MEMBER LEITCH: What would failure have
11	been in this test, excessive strain rate? Or what
12	would you have construed as failure?
13	MR. BASU: Obviously, one definition would
14	be it pops open, but what we were obviously looking
15	at.
16	MEMBER LEITCH: Obviously that, but I was
17	wondering if you had a lower threshold of failure?
18	MR. BASU: Yes. If it had gone from a
19	secondary creep or the secondary creep regime to
20	tertiary creep regime would the strain have gone
21	substantially up. We would consider that to be at
22	least close to the failure.
23	And that we don't have.
24	CHAIRMAN POWERS: It sure looks like No.
25	4 was getting close.

1 MR. BASU: I can give you the bottom line. 2 Everything is fine and dandy and nothing happened with 3 these rods, but let me just go through a couple of 4 slides here to show you in terms of plots, some 5 interesting observations. 6 Returning to 400 C. at 198 MPa and 380 7 degrees C., so that's a matter of 20 degrees difference. 8 There was obviously a significant 9 difference in hoop strain all the time. Likewise, if 10 you go from 190 MPa to 220 MPa, we saw 11 significant difference in hoop strain and again, 12 nothing unexpected. This is what you would expect by 13 increasing the temperature or by increasing the 14 stress. 15 What was obviously not obvious to us is 16 that by increasing 20 degree temperature, that you're 17 going to see that much difference in hoop strain. 18 Then, if combine, you make some 19 combinations of stress and temperature, you 20 actually get the same kind of strain for both 21 combinations and in this case we're showing that 380 22 to 220 MPa is very similar to 400 degrees C., 190 MPa, 23 similar kind of strain. Again, what that tells us is that in the 24

laboratory environment we can actually keep one of the

two parameters, the temperature or stress, very representative of what would be the beginning of life of dry storage condition and then we can artificially increase or decrease the other parameter, but then we can come back to analytically to what we would expect to see in terms of the real parametric changes.

## Where am I?

Okay, this one is what Dana, you asked me and I tried to explain what we did is we basically ran that 190 degree and -- I'm sorry, 190 degree, what is that? 400 degree, 190 MPa test up to 1870 hours or so. It was not much happening in terms of strain accumulation. Then we jacked the stress up to 250 MPa and you can see that it is going really fast. But still in the steady state regime.

This plot, only to show that the average strain that we have been talking about all along is actually pretty close to the outer diameter strain that we also measured.

## Okay?

What are the conclusions? Significant creep, residual creep strain is demonstrated, even up to 15 years of dry storage. So one implication is that you can go on for another extended period of time without accumulating a lot of strain and realize, of

course, the temperatures are now actually up to 15 years, even much lower than beginning of light temperatures and the pressure also.

The creep ratio, strong temperature and stress dependency and the regime tested, we haven't tested tertiary regime. We haven't been able to take anything to tertiary regime as yet, but in the steady-state regime that's the dependence.

Now coming back to the, let's see, ah, No. 6 and 7 which have not been done yet. It's not complete yet. What we do want to do 400 C., and different stress level because our Spent Fuel Project Office is in the midst of revising the interim Staff Guidance 11. The original guidance has a temperature limit of, I believe, 380 C. or 360 C. and they're looking into the prospect of actually describing a 400 C. temperature instead of 360 or 380. There's no one here now from the Spent Fuel Project Office, so I'm not sure if I'm -- I think I'm representing them okay in terms of their intent, but they can verify that.

CHAIRMAN POWERS: It seems to me that you're generating a data base with the sufficiency which you could accommodate a licensee coming in and saying well, I want to run it at 400 or some range of temperatures and you can say well, that's okay.

1 MR. BASU: Yes, that's what this data is 2 showing at the moment. Of course, the other thing 3 with the 400 degrees is the fact that at 400 degrees 4 as we have seen from the vacuum-drying process that 5 you have some hydrogen that will go into solution and then they will again reprecipitate and whether or not 6 7 in that process there is any reorientation. We haven't see, of course, at Surry, but with Robinson 8 9 rod campaign that's probably another story. 10 So let's see, have I gone through that? 11 There it is. High burnup. In essence, it is very 12 similar to the Surry campaign that we had concluded. 13 We're going t.he fuel and cladding to do 14 characterization. We have already started that. We're going to do isotopic analysis. Ralph alluded to 15 16 that. We have performed annealing tests to again see 17 whether or not there have been some annealing that 18 took place already. 19 We have put a lid test specimen in the 20 furnace for tunnel creep test and that's in July, back 21 in July. 22 CHAIRMAN POWERS: You have a ways to go 23 yet. MR. BENNETT: That's right, a ways to go, 24 25 that's right. We'll do some mechanical properties

285 The material is Robinson, as I said, and 67 1 test. gigawatt day, burnup, 2.9 percent enrichment. 2 oxide thickness is between 60 micron and 110 microns. 3 The hydrogen content is anywhere from 600 to 750, 800, 4 5 perhaps. status the analysis isotopic is 6 characterization in 7 analysis in progress, the progress. Annealing test completed and I'll show you 8 some results. 9 Creep test matrix developed. Now, when we 10 were about to start Surry creep testing, we came up 11 with a creep test matrix. We have done a kind of peer 12 In fact, we had two peer reviews of review of that. 13 that test matrix in terms of its progression to come 14 up with the final test matrix and a lot of that 15 actually depended on what we predict as going to be 16 some model, some creep's trend based on 17 the

> And this is what we have to do in terms of the development of the Robinson test metrics. The lead tests started and the mechanical testing plan.

> correlation and what we actually observe as we started

this creep testing and then we changed or modified our

What am I showing here? Oh, this is the annealing test results which we completed and all this

course.

18

19

20

21

22

23

24

table shows that there is irradiated rods with 600 PPM hydrogen. The peak DPH number that we came up with is 252 which is -- what it says is it's very close to not having any annealing. That's what it says.

What is the testing strategy? We're going to conduct two lead tests, two duplicate Surry tests which show that everything is in order. We're going to -- one has started, as I said already. Then we're going to establish test methods based on the lead test results to see whether or not we are getting the kind of strain that the models are predicting and we're going to emphasize 400 degrees. I mentioned the reason earlier and we're going to duplicate the testing technique.

slide giving you the last Ι preliminary creep test matrix. It doesn't really do It does give you the temperature, justice here. indicating that we are focusing on 400 degrees C. and of course a couple of tests around 400 degrees C. are focusing on stress where we think that we are going to have reasonable and measurable creep strain. We don't know about the duration that we're going to subject this test to. That will be determined based on the lead test results and of course we can predict creep trend based on the current model, current

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

1	correlation. We just don't know how good the
2	prediction is going to be so this is still to be
3	computed.
4	And that's about it.
5	MEMBER LEITCH: The Robinson rods have not
6	been in a cast for 15 years.
7	MR. BASU: That is correct. That is
8	correct.
9	MEMBER LEITCH: This is like a baseline?
10	MR. BASU: Well, if you look at the if
11	you are actually wondering what the creep test matrix
12	in this case, the beginning of life temperature is
13	probably going to be similar to what I have shown.
14	That's not going to change.
15	Now when we took these Surry rods out,
16	after 40 years in dry storage or 50 years, the
17	temperature was something on the order of 150 degrees,
18	rather than 360 degrees, but our tests are based on
19	beginning of life.
20	CHAIRMAN POWERS: Now the Surry specimens
21	have been in dry cask storage for 15 years. So
22	they've seen a fair amount of creep already.
23	MR. BASU: You saw the amount of creep
24	they saw which is less .1 percent.
25	CHAIRMAN POWERS: Because of the stress,

presumably. 1 You've seen the logarithmic creep --2 MR. BASU: Yes, of course. 3 Lower stress than what the trends that we tested at, 4 Absolutely. 5 RZ: I heard Carl Papariello lecturing and 6 7 he was outright eloquent and what he said was look, this stuff is going to go into an ISFI, it's going to 8 be there for some indeterminate number of decades and 9 some days some future generation of engineers is going 10 to open this thing up and take this stuff out, without 11 saying what one might do with it at that point. 12 And he didn't want it to fall apart on 13 You're going to get some hook or clamp or 14 them. 15 something and pull it out and it shouldn't fall apart. We shouldn't leave a problem for another generation 16 and he said it better than I just did, but that's the 17 goal. And I think what we're generating is some sound 18 data putting this on a data base. Things are okay. 19 20 And there's nothing wrong with a good news story. MR. BASU: May I have just the last word? 21 CHAIRMAN POWERS: Sure. 22 This program was co-sponsored 23 MR. BASU: by EPRI and DOE, Office of Civilian and Radioactive 24 Waste Management. So this is a joint program and I'd 25

1	like to acknowledge EPRI. EPRI representatives are
2	here. DOE is not here, but DOE was an equal partner
3	in this program.
4	CHAIRMAN POWERS: Golly darn. Thank you.
5	Are there any other questions for Dr. Basu?
6	I'm starting to lose the ability to talk
7	and I haven't even been speaking. Sud's comment
8	prompts me to ask did I mention that the LOCA work at
9	Argonne was done in cooperation with EPRI.
10	CHAIRMAN POWERS: You did.
11	CHAIRMAN POWERS: Good.
12	MEMBER ROSEN: And that cooperation with
13	the utilities who fund EPRI. EPRI has no money.
14	CHAIRMAN POWERS: I take it that this is
15	a paid political announcement here.
16	MEMBER ROSEN: The preceding was a factual
17	
18	CHAIRMAN POWERS: Statement of fact.
19	RZ: We have a full Committee summary and
20	if you have some direction.
21	CHAIRMAN POWERS: Give me time. We'll get
22	to that. First of all, I'd like to thank all the
23	speakers for an extremely informative sessions,
24	excellent presentations on the part of all and it
25	filled the Committee with information.

It comes time now for the Committee to work and I had said that the Committee should think about two questions. One is what should be presented to the full ACRS and I will cast out a preliminary agenda.

Our focus in discussions with the Committee is, in fact, on the RES research program and we have in our second question a debate on what we actually want the ACRS as a whole to do here, but I would suggest that any presentation to the Commission focus heavily on that RES program as it stands now because that's the issue that we confront right now.

I would suggest the following that we -that I begin with an opening summary of the general
issue in which I can give a thumbnail sketch of the
presentations that EPRI made in this. It is not
because I didn't think the EPRI work is excellent.
It's that the ACRS as a whole does not have to
confront that particular issue until NRR comes back
with their SER on the issue.

If we tell them all this wonderful stuff that we heard today at the meeting, they will simply forget. By the time the SER comes, because as we heard from Undine, there's a fairly deliberate program to review that material underway. And I think when

Now this is my proposal to the Committee 1 and you guys are free to augment this. And then it 2 seems to me that following Ralph's program might be 3 the appropriate time for Undine to give us 4 description of what you're planning to do on the 5 review of the EPRI work. I mean I would -- you had a 6 fairly succinct presentation of that plan that you 7 presented here and I think that's about an appropriate 8 level of detail which I have to give a little bit more 9 introduction on the issue, just so they can put it in 10 the context. 11 That's my proposal. 12 MEMBER FORD: You will give a synopsis of 13 14 the EPRI program to start? will start the CHAIRMAN POWERS: Ι 15 Committee off with getting them back up to speed on 16 what the overall issue is and in the course of that, 17 I will -- in connection with the RIA, give a capsule 18 summary of the approach that you outlined in the 19 analysis, the ductility approach that you've taken and 20 the separation you have between the coolability and 21 the rupture limits there. Does that sound fair? 22 Sounds fair. MS. YANG: 23 So I'll take a little CHAIRMAN POWERS: 24 more time in the introduction than is common, but I 25

that evaluation report comes from NRR that would be the appropriate time for EPRI to present the material to the full Committee and perhaps even remind this subcommittee of all the material because I'm sure there would be more and better understanding that will come along at that time.

So it's just that the press of things will mean that the ACRS will simply forget and so there's no real need to do that whereas they're focused very much on the research program. Then we would ask Ralph who taking as a springboard his opening presentation perhaps augmenting with and to us presentations of some of the new results you've gotten in the area of LOCA, some of your new thinking about how to approach the RIA and some of your thinking about the ATWS, you can take a bulk of the time to get the Committee up to date on where you stand in your research program.

I think in the course of that I forgot to mention this business that you're reworking the program plan. I know you're not in a position to say what that rework program plan is, but you're going to have to mention that that's going on and give us some hint on when we will know when the new program plan becomes available.

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

think it's approach to do so because the Committee 1 loses track of where this issue is and in addition, as 2 Peter will be glad to tell you, there are several new 3 Members who haven't had the benefit of all the history 4 in this program and what not. So I'll take a little 5 6 more time to begin. MEMBER FORD: Rosa, did you want to say 7 something else? 8 I would just -- maybe some 9 MS. YANG: There's an inconsistency in what we 10 clarification. proposed and what Ralph talked about. From what you 11 just said, Dana, you don't think tomorrow is the place 12 to acknowledge that inconsistency. 13 CHAIRMAN POWERS: That's right. 14 MS. YANG: I agree with that. 15 CHAIRMAN POWERS: It think it's going to 16 be difficult for me to avoid saying there's 17 inconsistency, but I don't want to try to highlight 18 that right now. I want to say you guys have done a 19 detailed analysis and an approach on this problem, 20 given an outline of what it is that you've done and 21 I'll say at the end of the day's presentation, Undine 22 will talk about what NRR is doing to review that. 23 I don't think Ralph wants to contest what you said 24

right now. You certainly didn't today. He simply has

a different approach and he gets a little more time to 1 2 outline his approach, but his is still a work in 3 progress and that's what the Committee needs to know 4 about. 5 We're in the business of advising the Commission on the viability of this and I think the 6 7 time to try to get a common view on that is when we 8 have the NRR review, the work. 9 SIEBER: They're not necessarily MR. 10 inconsistent. 11 CHAIRMAN POWERS: They're not necessarily 12 inconsistent. 13 MS. YANG: The only thing I want to point 14 out, I agree they're not necessarily inconsistent in 15 many aspects of it, but one of the aspects which is 16 extremely critical to the industry which 17 separation of coolability and fuel failure 18 because for fuel failure you calculate the dose and we 19 all know how to do that and we have done that. But 20 coolability is the safety limit and that's the most 21 important limit. And I just don't think there's any 22 discussion yet. 23 Our comment on what Ralph proposed, and we 24 have gone through a lot of discussion regarding our

coolability limit, so I'm a little bit concerned about

to bring that issue too much forward in the limited 1 time because that point is of major importance to us. 2 I think the fact that CHAIRMAN POWERS: 3 it's a major point, it's unavoidable for me saying 4 that to the Committee. I just don't think I can avoid 5 saying that, but I don't think I want to resolve it 6 7 here. MEMBER ROSEN: I don't think the Committee 8 will have any interest in trying to resolve it either. 9 CHAIRMAN POWERS: They're going to draw a 10 blank. 11 That's right, but it will MEMBER ROSEN: 12 be necessary for you to say this is difference in 13 approaches and that the significant impact of that 14 difference. 15 MR. SIEBER: I don't think it's resolvable 16 in the time that we have, number one because you're 17 going to have to get into a lot of detail to do that 18 and I don't think anybody is really prepared, maybe 19 EPRI is, but I don't think the rest of us are prepared 20 to deal with that issue to finality at this point. 21 MS. YANG: No. 22 CHAIRMAN POWERS: If we're going to get --23 some time in December, we're going to know a schedule 24 of when NRR is going to have an in-depth review and I 25

think it's once that review comes forward that we're in a position to discuss the nitty-gritty of those issues and right now we're really working on the design of the research program subject, of course, to whatever comes out of this revised program plan that we've done not too much about, but I mean I think it will still have the same elements that we're going to hear about, RIA, LOCA and ATWS. There may be a different emphasis across that board and of course I left out the spent fuel work, but that seems to be progressing along at a nominal pace. It would be really good if we could 12

declare something done. And just programmatically, if had my druthers, I would finish the 1998 program and work out a new program and call it a new program. go into advanced field -- and then I could say this would be a great value to us.

I think that's great. CHAIRMAN POWERS: You're stuck with the fact that these tests with irradiated fuel don't conform well to management's And I think we have to live with that. schedule. think the Committee's interest in knowing what's going on -- by the way, Ralph, the Committee will be very interested in the CABRI test matrix. You didn't put it up in your presentation, but I think Rosa had a

1

2

3

4

5

6

7

8

9

10

11

13

14

15

16

17

18

19

20

21

22

23

24

very nice slide in her presentation, something akin to that.

Perhaps when you're discussing what's going to be accomplished at the end of 2003 with an analysis in 2004, and then you can show the follow-on confirmatory tests and what not. I think the Committee is very interested in this because we did years ago write a letter endorsing that cooperative agreement and like to know where they're coming along.

My proposal. Now the second question is whether we should write a letter here and at this point I'll turn to Peter and say you have an alternative to writing a letter on the research program at the end of this meeting.

MEMBER FORD: Yes. You started off the meeting, Dana, by saying that there would be a letter because the assumption was that this particular topic would not be in the scope of the ACRS report on the RES plan for advanced reactors.

In writing out the scope of that report,

I put it that we really should be looking at where we
will be in 20 years time in terms of the reactor
fleet. My guess is we'll have our current reactor
fleet upgraded, obviously, and license renewed. In
all likelihood from the risk perspective, advanced

light water reactors coming potentially on line and maybe we might have a gas cool reactor. That's a real 2 3 stretch in my view. But regardless in the time period that we 4 have in 2003-2004 working period, if we just look at 5 the time lines, you've got a huge gap. You've got an 6 The research that you guys have got to do 7 with respect to some of the advanced light water 8 reactors and especially gas reactors, and then the 9 industry has to make some commercial decisions. 10 So in that time period, the fuels, for 11 instance, high burnup fuels, MOX fuels have kind of 12 limited my experience with this, but there must be 13 some areas which are on-going in our current programs 14 and the advanced reactor program which have to be done 15 on a priority basis right now as it impacts where we 16 will be in 2020, 2025. 17 MR. SIEBER: Well --18 Just to finish up, Jack, I MEMBER FORD: 19 think that's why some of this project that we talk 20 about, high burnup fuels, is relevant to the advanced 21 reactor coolant. That's my suggestion being that some 22 of this work is appropriate for the ACRS report on the 23 advanced reactor program. 24

MR. SIEBER:

Maybe I could comment a

299 little bit on a couple of things. If I looked at 1 future reactors, it seems to me the work is being done 2 now for the current fleet is applicable to advanced 3 light water reactors. This appears that way to me. 4 Gas cooled reactors is not clear to me 5 whether they'll be deployed or not and if I look at 6 the roadmap for June 4, deployment is 25 years in 7 advance and so starting something next fiscal year for 8

any of those concepts is probably premature.

On the other hand, I think that we have to recognize that they're out there and be prepared at least with some conceptual plans as to what research should be about to put our arms around any one of those concepts.

I'd like to get back to the issue of what gets said to the Committee. One of the artifacts that has been laying around for several months is RepNa-1 test data which caused some excitement and I think it would be worthwhile to say a sentence or two or at least consider saying it that the data that came out of that was, isn't considered to be an outlier and I think that there is a firmer basis to establish conservative limits without saying that this is a valid data point.

And I think you can take it or leave it,

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

which is up in the range of 60, 80, 100 calories per 1 gram or maybe higher than that, the pulse widths 2 should be around 10 calories per gram because in a PWR 3 you just could not, it --4 CHAIRMAN POWERS: You mean 10 millisecond 5 pulses? 6 Did MR. MEYER: Ten millisecond pulses. 7 I misspeak? 8 CHAIRMAN POWERS: You said 10 calories per 9 10 gram. MR. MEYER: Ten millisecond. It's getting 11 12 late. So this is a point that we've been making 13 over and over again in our discussions with the 14 industry and with the CABRI Technical Advisory Group 15 as they plan future tests, because they continue to 16 plan these tests with a 30 millisecond width. 17 Brookhaven has also looked at 18 dilution events to look at the power level, the pulse 19 widths and I have a few of those slides. I think I'll 20 sort of rush through them in order to save a little 21 I won't skip them all together, more time for Sud. 22 but two cases are illustrated here, one with pumps on 23 them and one with pumps off. This is the power and 24 boron concentration and it shows these spikes. 25

10

11

12

13

14

15

16 17

18

19

20

21

22 23

24

25

This event is very reminiscent of the BWR power oscillations from the worm's eye view, from the It looks very similar. And what fuel point of view. you see is you see peak fuel enthalpy from the first pulse is very low. So you can quickly get in a little bit of fuel enthalpy and then you can get in more which is also the case in the BWR oscillations, but it happens more slowly and during that time you get heat transfer and the cladding heats up.

The cladding is then able to take it to expand, to deform and so it appears in this case to me just at first blush as it did to the PIRT members look at ATWS that probably the PCMI is not going to be the big challenge for the fuel, but rather the temperature excursion.

How do you get a SIEBER: MR. dilution, a boron dilution that fast? What's the phenomenon in the plant that would take you from --

MR. EL-ZEFTAWY: Actually this was GSI 185 We spent a whole day with and it's being reviewed. Subcommittee, what's Thermal-Hydraulics but the postulated is that you have a small break LOCA and in a BNW plant. You've effectively distilled water. You now have a slug of unborated water in the -- down by the loop seal and then you do one of two things. You

through natural phenomena, natural 1 either restarts which is slower, or the operators start the 2 And their procedures tell them not to, but 3 that's how you could get these sort of events. 4 But that's well beyond the 5 MR. SIEBER: design basis, right? 6 7 MR. EL-ZEFTAWY: It looks like something 8 that almost happened. I'm sorry. There's an error in this 9 MR. MEYER: This is peak fuel enthalpy, peak 10 This is the other case, natural 11 enthalpy. circulation, the power and boron. Well, I guess it's 12 just the power curves and the enthalpy curves, labeled 13 14 correctly. Now moving on from boron dilution to BWR 15 rod drop, we've not focused much on BWR rod drop 16 because in our risk perspective, we thought that the 17 power oscillations were more important to look at than 18 The rod drop has a lower probability 19 the rod drop. than the rod ejection in the PWR, so we haven't spent 20 much time on it and we still haven't spent much time 21 on it. Brookhaven had done some earlier calculations. 22 They went back and had a look and it appears, I will 23 just say it appears that the pulse width for the 24

boiling water reactors are indeed broader than for the

but we made a fuss about it at one time and I'm sure that it will come to others' minds if it comes to mind.

CHAIRMAN POWERS: Yes. I mean, it seems to me that in the EPRI presentation that Rosa made there was a discussion of rather elaborate efforts that they'd been going to try to understand this test. I would certainly bring that up in a summary presentation.

MR. SIEBER: Great.

CHAIRMAN POWERS: And I would say their conclusion is that this is probably an outlier or difficult to explain.

Ralph, in your presentation you might want to think about putting in just a slide or two, say a slide or at least a line on a slide that outlines Hee's point, Hee Chung's point and you indicated that you, too, are prepared to say that this is an outlier, that doesn't have to fit all the correlations here and I agree with Jack. There are two things that have impressed me today as take home lessons. One is there is a burnup dependence to the enthalpy the fuel will take and that there seems to be an agreement that RepNa-1 is a peculiar test. That seems to be a point of agreement that is significant to my mind.

MEMBER ROSEN: I have one other thing that 1 I think I can take away and that is the information in 2 3 the dry cask storage. I think that is something that should be mentioned. 4 5 CHAIRMAN POWERS: Yes, I've left that out. 6 Sud, did you want to say something to the Committee? 7 MR. BASU: I want to remind you that this 8 was the medium burnup work. I think all of your other 9 presentations were high burnup, so I did not know how 10 you plan to -- I don't know how you plan to couch the 11 medium burnup work, but I think there is a value to 12 this work in the sense that we are going to follow the same procedure, same testing methods and the campaign 13 would be pretty much the same. So we are going to 14 15 generate some high burnup data soon. But notwithstanding the 16 MEMBER ROSEN: 17 fact that you've got to go on and do high burnup work. I think the results you presented today are valuable 18 19 for the Committee to know that there has been an 20 organized look at some fairly long stored medium burnup fuel and that the results are nominal. 21 22 MR. SIEBER: Just make the cask 10 inches 23 longer --CHAIRMAN POWERS: 24 The Committee, the Planning and Procedures Committee has only given us an 25

1	hour and 25 minutes and I'm trying to avoid having
2	people racing up here like scared deer
3	MR. BASU: Dana, I don't have to make a
4	statement in the meeting.
5	CHAIRMAN POWERS: Could you perhaps arm
6	Ralph with two or three vu-graphs so that he could
7	just give a capsule statement on the existence of the
8	work and indicate that it's going on.
9	MR. MEYER: I have those already captured.
10	CHAIRMAN POWERS: Maybe that will be
11	useful to begin because I agree very much with Steven.
12	That's not a usual thing for the two of us to agree.
13	MEMBER ROSEN: I promise not to do it
14	again.
15	(Laughter.)
16	CHAIRMAN POWERS: And what you're
17	essentially coming back to so far okay, Ralph, I
18	can count on you capturing that because I agree with
19	Steve, that that's a significant point.
20	Are there other comments to be made?
21	MEMBER FORD: I still didn't hear a
22	conclusion about whether we have a letter or not.
23	CHAIRMAN POWERS: Here's what I would
24	propose the Members of the Subcommittee to do. I'll
25	ask you to think about it tonight and give me some

advice tomorrow on whether we'll write a letter and regardless of what your position, if a letter is to be written, any points you think are to appear in it, what not.

My tendency is to go ahead and write a letter on this program, because I think it has some visibility with the Commission. I think there's been

Right now, I think those comments are fairly benign in the sense that they say progress has been made and is being made and stay tuned. I don't think I have outstanding advice to give the researchers on how to do their work better. I don't think that there are any major changes in the direction here, but I have a tendency to think that this has -- there's enough money invested in this program that has enough visibility because it's a

some substantial investment in it. I think that it

If it seems appropriate to add more material into the overall research program, I think we can do that.

highly cooperative international program that we ought

to tell the Commission something about it, so that

they're aware of it. That's my general feeling.

MEMBER LEITCH: Just a couple of points

## NEAL R. GROSS COURT REPORTERS AND TRANSCRIBERS 1323 RHODE ISLAND AVE., N W WASHINGTON, D C. 20005-3701

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

merits comment.

that I had and I think in your synopsis of the EPRI 1 presentation, certainly discuss that there's a burnup 2 dependent failure limit. I guess what I think I heard 3 today is that it may actually be more correct to say 4 there's an oxide film thickness dependent failure 5 limit, but burnup is more easily measurable circuit 6 for that perhaps. 7 Well, I think it's a CHAIRMAN POWERS: 8 ductility argument that's being advanced and in truth, 9 I think that's why Jack sees there's not a great 10 controversy between the two because I see Ralph 11 talking about things that smack of ductility here as 12 13 well. The other thing I heard MEMBER LEITCH: 14 There was an allusion to a that was interesting. 15 future presentation on the Robust Fuel Program. Ι 16 hope that doesn't get lost in the shuffle some place. 17 I think we need to --18 CHAIRMAN POWERS: Rosa and I have agreed 19 that some time after the first of the year, but we'll 20 talk on the phone. 21 MEMBER LEITCH: Okay. 22 CHAIRMAN POWERS: There are two things, it 23 seems to me, I think there's a lot in that program and 24 so I'm wondering if it shouldn't have a subcommittee 25

1	meeting to hear all about it, some time immediately
2	before a full Committee and give the full Committee a
3	synoptic picture of that whole program.
4	MEMBER ROSEN: I think it merits a
5	subcommittee meeting all by itself.
6	CHAIRMAN POWERS: It's a big program
7	that's been going on and I know Rosa is not very
8	enthusiastic about it and never thinks very much about
9	it, but I will implore to come give us a few words.
10	MEMBER ROSEN: She also knows if she comes
11	to speak about the Robust Fuel Program for a whole day
12	she can bring some supporting cast. She doesn't have
13	to do it all by herself.
14	CHAIRMAN POWERS: I was going to see her
15	do it by herself.
16	. · MEMBER LEITCH: And just one other
17	comment, maybe it's more in the form of a question for
18	Ralph, your second slide was titled "Original List of
19	Issues." And I'm not sure of the research plan that
20	you're working on. Are there different issues or are
21	we just further refining the resolution of these
22	original issues? It's not clear to me whether they're
23	new issues related to high burnup fuel that are going
24	to surface.
25	MR. MEYER: I don't think they are new

1	issues of that nature. There are, of course questions
2	about alloy effects for M5 and ZIRLO which are not
3	addressed in the current wrap up of the old issues,
4	but which are to some extent being planned in the
5	program. And those haven't been laid out in terms of
6	just what are we going to do and what are the
7	schedules for that. So that will constitute part of
8	the new program plan, but not necessarily represent
9	any new issues.
LO	MEMBER LEITCH: So there might be an
11	additional issue or sub-issue related to cladding
12	materials?
13	MR. MEYER: Yes, related to the cladding
14	materials. Yes.
15	CHAIRMAN POWERS: Well, I think we just
16	have to stay tuned for this new program plan. I got
17	the impression in the opening remarks that this is
18	very much a work in progress and maybe the progress
19	has just been initiated or something like that.
20	MR. WERMIEL: It hasn't just been
21	initiated, Dr. Powers, but it is a work in progress.
22	There's been discussion between the two
23	offices, actually three offices, because it is going
24	to include the NMSS piece as well and those
25	discussions have been going on for several months at

## (Laughter.)

And with that I think we can adjourn this subcommittee meeting with the imposition that all the Members should think about the points that should be raised in the letter on the research program and your advice on whether it's appropriate to write one or not. With that, I'll adjourn this meeting of the subcommittee.

(Whereupon, at 5:32 p.m., the meeting was concluded.)

least, but we still have certain things that we're 1 trying to clarify and clear up. 2 I think it's CHAIRMAN POWERS: 3 premature for the ACRS to try to inject itself into 4 this debate. 5 MR. WERMIEL: I think so, too. 6 CHAIRMAN POWERS: Any other comments 7 people would like to make? 8 Again, I really want to emphasize to all 9 the speakers that the presentations were excellent. 10 They were filled with information and I envy you all. 11 It looks like fun work and challenging work to sort 12 these things about. 13 I have to admit that I was just stunned at 14 the amount of work that must have been done, the EPRI 15 work because Robbie would get up there and say well, 16 here's a point and we did this with multiple computer 17 code calculations and things like that and he had 85 18 So I know there's a huge points like that, data. 19 Similarly, Ralph, you and amount of work there. 20 Harold, I know that each of your data points is 21 obtained with a great deal of pain and frustrations 22 and problems, so I very much appreciate you sharing 23 with us and Undine, I wish you well on your review 24 25 plan.

## CERTIFICATE

This is to certify that the attached proceedings before the United States Nuclear Regulatory Commission in the matter of:

Name of Proceeding: Advisory Committee on

Reactor Safeguards Reactor

Fuels Subcommittee

Docket Number:

A/N

Location:

Rockville, Maryland

were held as herein appears, and that this is the original transcript thereof for the file of the United States Nuclear Regulatory Commission taken by me and, thereafter reduced to typewriting by me or under the direction of the court reporting company, and that the transcript is a true and accurate record of the foregoing proceedings.

Velvera Davis

Official Reporter

Neal R. Gross & Co., Inc.